

FEATURES
SOURCE

Query Match	13.08;	Score 120.4;	DB 9;	Length 149940;
Best Local Similarity	61.58;	Pred. No. 5.1e-19;		
Matches 368;	Conservative	0;	Mismatches 156;	Indels 74;
				Gaps 8

08:25

[illegible]

REFERENCE
2 (bases 1 to 840)
Burglin, T. R., Matta, I. W., Newmeyer, D. D., Zeller, R. and de
Robertis, E. M.

CDS

polya-signal	634	639	174 g	129 t
BASE COUNT	222 a	121 c		
ORIGIN				

[illegible][illegible]

D**b**
149 GGCAAGCATGAGTTCACATAGTAGAAAT-----ADU1760841-1760849

[illegible]

us-09-844-864-16.rn1

[illegible]

Sequence 2057, Applicant
Patent No. 6090620
GENERAL INFORMATION:
APPLICANT: Fu, Ying-Hui
APPLICANT: Yu, Chang-En
APPLICANT: Oshima, Junko
APPLICANT: Mulligan, John T.
APPLICANT: Schellenberg, Gerald D.
TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO
TITLE OF INVENTION: WERNER'S SYNDROME
NUMBER OF SEQUENCES: 209
CORRESPONDENCE ADDRESS:
Fu and HARRY LLP

ADDRESSEE: Seed and Corn
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104-7092
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION NUMBER: US/08/781,891

APPLICATION NO: 27-DEC-1996
FILING DATE: 27-DEC-1996
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION: Ph.D., Carol
NAME: NO. 6090620tenburg
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 240052.419
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 209:
SEQUENCE CHARACTERISTICS:
LENGTH: 51259 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

Query Match Best Local Similarity	6.9%; 59.9%;
Matches 106; Conservative	<div>Pred. No.</div> 1.6e-06; <div>Mismatches</div> 71; <div>Indels</div> 0; <div>Gaps</div> 0;

RESULT 3
US-08-781-891-208/c
Application US/08781891

Sequence 2007-07-11
Patent No. 6090620
GENERAL INFORMATION:
APPLICANT: Fu, Ying-Hui
APPLICANT: Yu, Chang-En
APPLICANT: Oshima, Junko
APPLICANT: Mulligan, John T.
APPLICANT: Schellenberg, Gerald D.
TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO
TITLE OF INVENTION: WERNER'S SYNDROME
NUMBER OF SEQUENCES: 209

CORRESPONDENCE ADDRESS:
 ADDRESSEE: SEED and BERRY LLP
 6300 Columbia Center, 701 Fifth Avenue

STREET: 0000
CITY: Seattle
STATE: Washington

COUNTRY: USA
ZIP: 98104-7092

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;
; COMPUTER READABLE: Floppy disk
; MEDIUM TYPE: IBM PC compatible
; COMPUTER: IBM PC-DOS/MS-DOS
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
;
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/781,891
;

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FILED DATE: 27-DEC-1996
CLASSIFICATION: 800

ATTORNEY/AGENT INFORMATION:
NAME: NO. 6090620tenburg Ph.D., Car
REGISTRATION NUMBER: 39,317

REGISTRATION NUMBER: 240052.41
REFERENCE/DOCKET INFORMATION:
TELECOMMUNICATION INFORMATION:
(206) 632-4900

TELEPHONE: (206) 682-6031
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 208:

SEQUENCE CHARACTERISTICS:
LENGTH: 16442 base pairs
nucleic acid

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TYPE: nuclear
STRANDEDNESS: single
TOPOLOGY: linear

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US-08-781-891-208

Query Match	6.48;	Score 59.47;			
Best Local Similarity	57.68;	Pred. No. 1.1e-05;			
Matches 125; Conservative		0; Mismatches 91;	Indels 1;	Gaps 1;	

[illegible]

D_b 16369 CAGGACCAAGGAAGGCATCCTCGAGGAG 495
Q_y 436 GAGGAGAAGGAAGGACATCATGAGGATGAGGATGCACATATCTCTGAGGAG

GenCore version 5.1.3
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 24, 2003, 05:06:35 ; Search time 77 Seconds
(without alignments)
3680.121 Million cell updates/sec

Title: US-09-844-864-16
Perfect score: 924
Sequence: 1 cagccgcgtctctgcccgg.....tttgcgcgcgaagtatg 924

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: Issued_Patents_NA: *
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5: /cgn2_6/ptodata/2/lna/PCTUS_COMB.seq: *
6: /cgn2_6/ptodata/2/lna/backfiles1.seq: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	93.2	10.1	7218	1 US-08-232-463-14	Sequence 14, Appl
C 2	63.4	6.9	51259	3 US-08-781-891-209	Sequence 209, App
C 3	59.4	6.4	16442	3 US-08-781-891-208	Sequence 208, App
C 4	59.2	6.4	3489	2 US-08-728-323A-1	Sequence 1, Appli
C 5	59.2	6.4	3489	4 US-09-298-568-1	Sequence 1, Appli
C 6	59.2	6.4	32207	2 US-08-770-379-20	Sequence 20, Appl
C 7	59.2	6.4	32207	4 US-08-757-669A-20	Sequence 20, Appl
C 8	59.2	6.4	32207	4 US-09-230-371A-20	Sequence 20, Appl
C 9	57.6	6.2	3211	2 US-08-574-959A-8	Sequence 8, Appli
C 10	57.6	6.2	3211	4 US-09-357-014-8	Sequence 8, Appli
C 11	57.6	6.2	3901	2 US-08-574-959A-6	Sequence 6, Appli
C 12	57.6	6.2	3901	4 US-09-357-014-6	Sequence 6, Appli
C 13	56.6	6.1	2518	4 US-09-433-699-3	Sequence 3, Appli
C 14	55.2	6.0	2340	3 US-09-022-983-4	Sequence 4, Appli
C 15	55.2	6.0	2477	4 US-09-490-692-3	Sequence 3, Appli
C 16	54.4	5.9	9636	1 US-08-323-170B-1	Sequence 1, Appli
C 17	54.4	5.9	9636	4 US-08-954-441-1	Sequence 1, Appli
C 18	54.4	5.8	1236	2 US-08-741-134-5	Sequence 5, Appli
C 19	52.2	5.6	289	4 US-09-007-005-17	Sequence 17, Appl
C 20	52.2	5.6	289	4 US-09-244-796-17	Sequence 17, Appl
C 21	52.2	5.6	15378	3 US-08-785-420-1	Sequence 1, Appli
C 22	51.8	5.6	966	2 US-08-766-738-2	Sequence 2, Appli
C 23	51.8	5.6	966	4 US-09-262-610-2	Sequence 2, Appli
C 24	50.6	5.5	489	1 US-07-879-685B-3	Sequence 3, Appli
C 25	50.6	5.5	5183	1 US-08-459-568-3	Sequence 3, Appli
C 26	50.6	5.5	5183	2 US-08-399-411-3	Sequence 3, Appli
C 27	50.6	5.5	5868	3 US-08-516-859A-3	Sequence 3, Appli

28	50.6	5.5	5868	4 US-09-586-472-3	Sequence 3, Appli
29	50.6	5.5	5868	4 US-09-528-706-3	Sequence 3, Appli
C 30	50.4	5.5	1926	4 US-09-249-585A-4	Sequence 4, Appli
C 31	50.4	5.5	1931	2 US-09-130-114-2	Sequence 2, Appli
C 32	50	5.4	258	4 US-09-345-882-21	Sequence 21, Appli
C 33	50	5.4	6002	4 US-09-345-882-4	Sequence 4, Appli
C 34	50	5.4	162450	4 US-09-345-882-1	Sequence 1, Appli
C 35	49.4	5.3	3100	1 US-08-296-362-1	Sequence 1, Appli
C 36	49	5.3	2188	4 US-07-865-662F-10	Sequence 10, Appl
C 37	49	5.3	2188	4 US-08-374-219B-10	Sequence 2, Appli
C 38	48.8	5.3	1678	3 US-08-650-766-2	Sequence 2, Appli
C 39	48.8	5.3	1954	3 US-08-922-635-2	Sequence 2, Appli
C 40	48.8	5.3	2255	2 US-08-741-134-1	Sequence 1, Appli
C 41	48.8	5.3	3318	3 US-08-650-766-3	Sequence 3, Appli
C 42	48.8	5.3	3318	3 US-08-922-635-3	Sequence 3, Appli
C 43	48.8	5.3	3385	3 US-08-650-766-1	Sequence 1, Appli
C 44	48.8	5.3	3385	3 US-08-922-635-1	Sequence 1, Appli
C 45	48.8	5.3	15202	3 US-08-922-635-21	Sequence 21, Appl

ALIGNMENTS

RESULT 1
US-08-232-463-14/C
Sequence 14, Application US/08232463
Patent No. 5670367
GENERAL INFORMATION:
APPLICANT: DORNER, F.
APPLICANT: SCHEIFLINGER, F.
APPLICANT: FALKNER, F. G.
TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
NUMBER OF SEQUENCES: 52
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 1800 Diagonal Road, Suite 500
CITY: Alexandria
STATE: VA
COUNTRY: USA
ZIP: 22313-0299
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/232,463
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/07/935,313
FILING DATE:
APPLICATION NUMBER: EP 91 114 300.6
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30472/114 IMMU
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)836-9300
TELEFAX: (703)683-4109
TELEX: 899149
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 7218 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
CLONE: PT29pt-Fls
US-08-232-463-14
Query Match 10.1%; Score 93.2; DB 1; Length 7218;

Best Local Similarity 3.5%; Pred. No. 1.5e-14;
Matches 14; Conservative 257; Mismatches 125; Indels 0; Gaps 0;

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QY 364 TTCCTCAGTGGCCAGAACGTTATGAGCATCAGACCTAACCTGGGAGGAGGAGGAGAA 423
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Db 1441 TTGTACRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1382

QY 424 GAAGAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 483
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1381 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1322

QY 484 TCTCTGAGAGGAGCAAGCCCTGTCAACAAGTCAAAAGCTGTGCCCCAGAGCAGCGC 543
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Db 1321 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1262

QY 544 AGCGTGCTAAGAAAAAAGCTGGAAGAAAGAGAGGAATAGAGCCAGCGTTAGA 603
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Db 1261 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1202

QY 604 GACAAGAGCCCTGTGAAAAAGGCCAAGCCAGAGCCAGAGCCAGAGATTCAAG 663
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Db 1201 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1142

QY 664 AATGAGGAGGAGCCGCTTGGGGGACAGGTCGAAGTGCGCTTCCCTGGGCTGTGCTG 723
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Db 1141 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1082

QY 724 CAGGCACAGGGTGCCCTGTCCAGCCCTCCACCTG 759
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RESULT 2

US-08-781-891-209/c

; Sequence 209, Application US/08781891

; Patent No. 6090620

; GENERAL INFORMATION:

; APPLICANT: Fu, Ying-Hui

; APPLICANT: Yu, Chang-En

; APPLICANT: Oshima, Junko

; APPLICANT: Mulligan, John T.

; APPLICANT: Schellenberg, Gerald D.

; TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO

; TITLE OF INVENTION: WERNER'S SYNDROME

; NUMBER OF SEQUENCES: 209

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: SEED and BERRY LLP

; STREET: 6300 Columbia Center, 701 Fifth Avenue

; CITY: Seattle

; STATE: Washington

; COUNTRY: USA

; ZIP: 98104-7092

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/781,891

; FILING DATE: 27-DEC-1996

; CLASSIFICATION: 800

; ATTORNEY/AGENT INFORMATION:

; NAME: No. 6090620tenburg Ph.D., Carol

; REGISTRATION NUMBER: 39,317

; REFERENCE/DOCKET NUMBER: 240052.419

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (206) 622-4900

; TELEFAX: (206) 682-6031

; INFORMATION FOR SEQ ID NO: 209:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 51259 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

US-08-781-891-209

Query Match 6.4%; Score 63.4; DB 3; Length 51259;
Best Local Similarity 59.9%; Pred. No. 1.6e-06;
Matches 106; Conservative 0; Mismatches 71; Indels 0; Gaps 0;

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QY 408 GGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 467
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Db 257 GGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 198

QY 468 TGAGGATGCAGATATATCTCTGAGGAGGAGCAAGCCCTGTCAACAAGTCAAAAGCGTGT 527
    ||||| | | | | | | | | | | | | | | | | | | | | | | |
Db 197 AGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 138

QY 528 GCGCCAGAGCAGGCGAGCGTGCTAAGAAAAAAGCTGAAAAAAGAGAGAGGA 584
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Db 137 GAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 81
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RESULT 3

US-08-781-891-208/c

; Sequence 208, Application US/08781891

; Patent No. 6090620

; GENERAL INFORMATION:

; APPLICANT: Fu, Ying-Hui

; APPLICANT: Yu, Chang-En

; APPLICANT: Oshima, Junko

; APPLICANT: Mulligan, John T.

; APPLICANT: Schellenberg, Gerald D.

; TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO

; TITLE OF INVENTION: WERNER'S SYNDROME

; NUMBER OF SEQUENCES: 209

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: SEED and BERRY LLP

; STREET: 6300 Columbia Center, 701 Fifth Avenue

; CITY: Seattle

; STATE: Washington

; COUNTRY: USA

; ZIP: 98104-7092

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/781,891

; FILING DATE: 27-DEC-1996

; CLASSIFICATION: 800

; ATTORNEY/AGENT INFORMATION:

; NAME: No. 6090620tenburg Ph.D., Carol

; REGISTRATION NUMBER: 39,317

; REFERENCE/DOCKET NUMBER: 240052.419

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (206) 622-4900

; TELEFAX: (206) 682-6031

; INFORMATION FOR SEQ ID NO: 208:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 16442 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

US-08-781-891-208

Query Match 6.4%; Score 59.4; DB 3; Length 16442;

Best Local Similarity 57.6%; Pred. No. 1.1e-05;

Matches 125; Conservative 0; Mismatches 91; Indels 1; Gaps 1;

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QY 436 GAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 495
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1161 GGAGGACGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGATGACGA 1220

TITLE OF INVENTION: POLYPEPTIDES FROM KAPOSI'S SARCOMA-ASSOCIATED

TITLE OF INVENTION: HERPESVIRUS, DNA ENCODING SAME AND USES THEREOF

10

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QY	483 ATCTCTGGAGGAGCAAAGCCCTGTCCAACAAGTCAAAAAGCGTTGGTCCCAGAAACAGGC					542
Db	1471 AGAAGAAGAGGAGGAGGCCACAGATTCTGAAGAGGAGGAGCATCTGGAACAGATGCAGGA					1530
QY	543 GAGCGTGGCTAAGAAAAAACCTGGA AAAAGAAAGAGAGGAATAAGAGCCAGCGTTAG					602
Db	1531 GGTCACGAGAGGATGATGAAGAGGAGCAGAAAGGAAGAAGCACAGCTAAAGATGG					1590
QY	603 AGACAAGAGCCCTGTG					618
Db	1591 AGACAAGAGCCCCCATG					1606

Search completed: January 24, 2003, 18:46:33
Job time : 206 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 23, 2003, 14:06:24 ; Search time 45576 seconds

(without alignments)
590.025 Million cell updates/sec

Title: US-09-844-864-16

Perfect score: 924

Sequence: 1 cagccgcgtctctgtccg.....tttgcgcgcgaagcttatg 924

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database :

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2: gb_htg: *
3: gb_in: *
4: gb_om: *
5: gb_ov: *
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14: gb_vl: *
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31: em_htg_inv: *
32: em_htg_other: *
33: em_htg_mus: *
34: em_htg_pln: *
35: em_htg_rod: *
36: em_htg_mam: *
37: em_htg_vrt: *
38: em_sy: *
39: em_htgo_hum: *
40: em_htgo_mus: *
41: em_htgo_other: *

Pred. No. is the number of results predicted by chance to have a

Score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	883	95.6	1874	9 AK094267	AK094267 Homo sapi
2	337	36.5	423	6 AX321879	AX321879 Sequence
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4	262.4	28.4	166878	2 AC023288	AC023288 Homo sapi
5	262.4	28.4	181714	2 AC090442	AC090442 Homo sapi
6	262.4	28.4	184050	2 AC087819	AC087819 Homo sapi
7	262.4	28.4	184444	2 AC090471	AC090471 Homo sapi
8	171.6	18.6	86574	9 HS833B7	AL008637 Human DNA
9	122	13.2	135206	2 AC108486	AC108486 Homo sapi
10	120.4	13.0	149940	9 AC093557	AC093557 Homo sapi
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12	96.2	10.4	800	5 XLNUPLR	X04766 Xenopus lae
13	93.2	10.1	7218	6 I66494	I66494 Sequence 14
14	90.6	9.8	198421	2 AC125180	AC125180 Mus muscu
15	83.2	9.0	190604	2 AC122303	AC122303 Mus muscu
16	81.6	8.8	200052	2 AL772398	AL772398 Mus muscu
17	81.4	8.8	251076	2 AC097410	AC097410 Rattus no
18	81.2	8.8	2081	9 AK056804	AK056804 Homo sapi
19	79.8	8.6	43553	9 AL357372	AL357372 Human DNA
20	79	8.5	222259	2 AC113078	AC113078 Mus muscu
21	78.4	8.5	241666	2 AC118543	AC118543 Mus muscu
22	77.4	8.4	182740	2 AC115811	AC115811 Mus muscu
23	76.8	8.3	165077	10 AC084382	AC084382 Mus muscu
24	76.4	8.3	170032	2 AC116673	AC116673 Mus muscu
25	76.2	8.2	70391	2 AC110410	AC110410 Rattus no
26	75.8	8.2	144328	9 AC009695	AC009695 Homo sapi
27	75.8	8.2	163495	9 AC022716	AC022716 Homo sapi
28	75.8	8.2	170631	9 AC020751	AC020751 Homo sapi
29	75.8	8.2	170807	9 AC023812	AC023812 Homo sapi
30	75.8	8.2	179803	9 AC068314	AC068314 Homo sapi
31	75.4	8.2	168856	2 AF466883	AF466883 Mus muscu
32	75.4	8.2	169153	2 AC128501	AC128501 Rattus no
33	75.2	8.1	110000	2 AC098456_2	Continuation (3 of
34	75.2	8.1	303943	2 AC127311	AC127311 Mus muscu
35	74.8	8.1	177716	9 AC026188	AC026188 Homo sapi
36	74.6	8.1	152763	2 AC119715	AC119715 Rattus no
37	74.6	8.1	175456	2 AC115183	AC115183 Rattus no
38	74.4	8.1	147670	10 AC084020	AC084020 Mus muscu
39	74.4	8.1	205606	10 AL596204	AL596204 Mouse DNA
40	74	8.0	184370	2 AL589845	AL589845 Mus muscu
41	73.8	8.0	120044	2 AC129042	AC129042 Rattus no
42	73.8	8.0	208844	2 AC125070	AC125070 Mus muscu
43	73.8	8.0	215938	2 AC102646	AC102646 Mus muscu
44	73.6	8.0	93682	10 AL591805	AL591805 Mouse DNA
45	73.6	8.0	154733	2 AC131221	AC131221 Rattus no

ALIGNMENTS

RESULT 1
LOCUS AK094267
DEFINITION Homo sapiens cDNA FLJ36948 f1s, clone BRACE2005719, weakly similar to NUCLEOPLASMIN.
ACCESSION AK094267
VERSION AK094267.1 GI:21753294
KEYWORDS oligo capping; f1s (full insert sequence).
SOURCE Homo sapiens cerebellum cDNA to mRNA, clone_11b:BRACE2 clone:BRACE2005719.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1
AUTHORS Tanigami,A., Fujiwara,T., Shibahara,T., Goto,Y., Hirao,M.,

Shimizu, F., Wakebe, H., Ono, T., Hishigaki, H., Watanabe, T., Ozaki, K., Sugiyama, T., Irie, R., Otsuki, T., Sato, H., Wakamatsu, A., Ishii, S., Yamamoto, J., Isono, Y., Kawai-Hio, Y., Saito, K., Nishikawa, T., Kimura, K., Yamashita, H., Matsuo, K., Nakamura, Y., Sekine, M., Kikuchi, H., Kanda, K., Wagatsuma, M., Murakawa, K., Kanehori, K., Takahashi-Fujii, A., Oshima, A., Sugiyama, A., Kawakami, B., Suzuki, Y., Sugano, S., Nagahari, K., Masuho, Y., Nagai, K. and Isogai, T.
NEDO human cDNA sequencing project
Unpublished
2 (bases 1 to 1874)
Isogai, T. and Yamamoto, J.
Direct Submission
Submitted (04-JUL-2002) Takao Isogai, FLJ Project(HRI Team); 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba 292-0812, Japan
(E-mail: genomics@hri.co.jp, Tel: 81-438-52-3975, Fax: 81-438-52-3986)
NEDO human cDNA sequencing project supported by Ministry of Economy, Trade and Industry of Japan; cDNA full insert sequencing; Research Association for Biotechnology (RAB); cDNA library construction; Helix Research Institute (HRI) (supported by Japan Key Technology Center etc.); 5'- & 3'-end one pass sequencing; RAB, HRI, and Biotechnology Center, National Institute of Technology and Evaluation; clone selection for full insert sequencing; HRI and RAB; annotation: HRI and RAB.
Location/Qualifiers
1. 1874
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="BRACE2005719"
/tissue_type="cerebellum"
/clone_lib="BRACE2"
/note="cloning vector: pME18SFL3"
BASE COUNT 394 a 570 c 604 g 306 t
ORIGIN
Query Match 95.6%; Score 883; DB 9; Length 1874;
Best Local Similarity 100.0%; Pred. No. 3.2e-208;
Matches 883; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CAGCCCGCTTCTGCCCCGAGGACCATGAATCTCAGTAGCGCCAGTACGAGGAGAAAG 60
Db 992 CAGCCCGCTTCTGCCCCGAGGACCATGAATCTCAGTAGCGCCAGTACGAGGAGAAAG 1051
QY 61 GCAGTGACGACCGCTGCTGGGGCTGCGAGCTCAGTCAGAGAGGCGGACTTGGACCTTC 120
Db 1052 GCAGTGACGACCGCTGCTGGGGCTGCGAGCTCAGTCAGAGAGGCGGACTTGGACCTTC 1111
QY 121 AGACCCCGAGCTGAGGGGAGAGCAGAGCTGCGAGCTGTGCTTCATACGATTTGCTGGGG 180
Db 1112 AGACCCCGAGCTGAGGGGAGAGCAGAGCTGCGAGCTGTGCTTCATACGATTTGCTGGGG 1171
QY 181 GAGAAAGCCAAAGAGAGATGATCGCGTGAGATCTGCCCCAGCAAAACAGAGAGAC 240
Db 1172 GAGAAAGCCAAAGAGAGATGATCGCGTGAGATCTGCCCCAGCAAAACAGAGAGAC 1231
QY 241 AAGAAGATGACGCGGTCACCATTTGCTCAGTCCAGGCTCAGTCCCTCCATGCTCTCC 300
Db 1232 AAGAAGATGACGCGGTCACCATTTGCTCAGTCCAGGCTCAGTCCCTCCATGCTCTCC 1291
QY 301 ATGGTAGAGTGCAGCTTTCTCCCCAGTTACTTTCCAGCTCCGGGCTGGCTCAGAGACC 360
Db 1292 ATGGTAGAGTGCAGCTTTCTCCCCAGTTACTTTCCAGCTCCGGGCTGGCTCAGAGACC 1351
QY 361 GTGTTCTCAGTGGCCAGGAACGTTATGAGCATCAGACCTAACCTGGAGAGAGAGAG 420
Db 1352 GTGTTCTCAGTGGCCAGGAACGTTATGAGCATCAGACCTAACCTGGAGAGAGAGAG 1411
QY 421 GAAGAAGAGGGGAGAGAGAGAGAGAGAGATGATGAGGATGAGATGACAGAT 480
Db 1412 GAAGAAGAGGGGAGAGAGAGAGAGAGAGATGATGAGGATGAGATGACAGAT 1471
QY 481 ATATCTCTGAGAGAGCAAAAGCCCTGTCAAAACAAGTCAAAAGGCTGTGCCCCAGAGCAG 540
Db 1472 ATATCTCTGAGAGAGCAAAAGCCCTGTCAAAACAAGTCAAAAGGCTGTGCCCCAGAGCAG 1531

QY 541 GCGAGCGTGCGCTAGAGAAAAAAGCTGGAAAAAGAGAGAGAAATAGAGCCAGCGTT 600
Db 1532 GCGAGCGTGCGCTAGAGAAAAAAGCTGGAAAAAGAGAGAGAAATAGAGCCAGCGTT 1591
QY 601 AGAGACAAGAGCCCTGTGAAAAAGGCCAAAGCCACAGCCAGAGCCCAAGAGATTC 660
Db 1592 AGAGACAAGAGCCCTGTGAAAAAGGCCAAAGCCACAGCCAGAGCCCAAGAGATTC 1651
QY 661 AAGAAATGAGAGAGCCAGCCCTTGGGGGCGACGGTGCAAAAGTGGGCTTCCCTGGGCTGTG 720
Db 1652 AAGAAATGAGAGAGCCAGCCCTTGGGGGCGACGGTGCAAAAGTGGGCTTCCCTGGGCTGTG 1711
QY 721 CTGCAGGCACAGGCTGCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCAACAGGGGTG 780
Db 1712 CTGCAGGCACAGGCTGCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCAACAGGGGTG 1771
QY 781 TTGCGGGGGCAACATGAGAGCCCTCACCCTCACTCTCCACTTTCAGAGAGGCCCCAGT 840
Db 1772 TTGCGGGGGCAACATGAGAGCCCTCACCCTCACTCTCCACTTTCAGAGAGGCCCCAGT 1831
QY 841 GAAGAGCCCGACCTCGGGGTCAACATAAAGTTGCTGTGTCAGG 883
Db 1832 GAAGAGCCCGACCTCGGGGTCAACATAAAGTTGCTGTGTCAGG 1874
RESULT 2
AX321879 423 bp DNA Linear PAT 15-DEC-2001
LOCUS AX321879
DEFINITION Sequence 410 from Patent WO0172295.
ACCESSION AX321879
VERSION AX321879.1 GI:17906455
KEYWORDS human.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE 1
AUTHORS Reed, S.G., Lodes, M.J., Mohamath, R., Secrist, H., Benson, D.R., Indrias, C.Y., Henderson, R.A., Flinn, S.P., Algate, P.A., Elliott, M., Mannion, J. and Kalos, M.D.
TITLE Compositions and methods for the therapy and diagnosis of lung cancer
JOURNAL Patent: WO 0172295-A 410 04-OCT-2001;
CORIXA CORPORATION (US)
FEATURES
source 1. 423
/organism="Homo sapiens"
/db_xref="taxon:9606"
BASE COUNT 84 a 138 c 125 g 76 t
ORIGIN
Query Match 36.5%; Score 337; DB 6; Length 423;
Best Local Similarity 100.0%; Pred. No. 6.5e-73;
Matches 337; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CAGCCCGCTTCTGCCCCGAGGACCATGAATCTCAGTAGCGCCAGTACGAGGAGAAAG 60
Db 87 CAGCCCGCTTCTGCCCCGAGGACCATGAATCTCAGTAGCGCCAGTACGAGGAGAAAG 146
QY 61 GCAGTGACGACCGCTGCTGGGGCTGCGAGCTCAGTCAAGAGAGGCGGACTTGGACCTTC 120
Db 147 GCAGTGACGACCGCTGCTGGGGCTGCGAGCTCAGTCAAGAGAGGCGGACTTGGACCTTC 206
QY 121 AGACCCCGAGCTGAGGGGAGAGAGAGCTGAGGCTGTGCTCATACGATTTGCTGGGG 180
Db 207 AGACCCCGAGCTGAGGGGAGAGAGAGCTGAGGCTGTGCTCATACGATTTGCTGGGG 266
QY 181 GAGAAAGCCAAAGAGAGATGATCGCGTGAGATCTGCCCCAGCAAAACAGAGAGAC 240
Db 267 GAGAAAGCCAAAGAGAGATGATCGCGTGAGATCTGCCCCAGCAAAACAGAGAGAC 326
QY 241 AAGAAGATGACGCGGTCAACATTTGCTCAGTCCAGGCTCAGTCCCATGGTCTCC 300

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 166878)
AUTHORS Abola, A.P., Bruno, D., Conn, L., Dela Rosa, M., Faulkner, D.,
Federspiel, N., Glukhov, S., Hansen, N., Herman, Z.S., Hyman, R.,
Mao, J., Lam, B., Marathe, R., Miranda, M., Morehouse, A.J., Nguyen, M.,
Oefner, P., Palm, C.J., Ramirez, D., Southwick, A.M., Wilhelmy, J.,
Yu, S. and Davis, R.W.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 166878)
AUTHORS Bruno, D., Conn, L., Dela Rosa, M., Faulkner, D., Federspiel, N.,
Glukhov, S., Hansen, N., Hyman, R., Mao, J., Marathe, R.,
Morehouse, A.J., Oefner, P., Palm, C.J., Ramirez, D., Wilhelmy, J.,
Yu, S. and Davis, R.W.
TITLE Direct Submission
JOURNAL Submitted (11-FEB-2000) DNA Sequencing and Technology Center,
Stanford University, 855 California Avenue, Palo Alto, CA 94304,
USA
COMMENT On Jul 19, 2000 this sequence version replaced gi:8980905.
----- Genome Center
Center: Stanford DNA Sequencing and Technology Development
Center
Center code: SDSRDC
Web site: http://sequence-www.stanford.edu/group/human/
Contact: hum-info@sequence.stanford.edu
----- Project Information
Center project name: 836
Center clone name: RP11-67H12
----- Summary Statistics
Sequencing Vector: M13mp18; X02513
Chemistry: Dye-primer; 0% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 151238 bases at least Q40
Consensus quality: 155909 bases at least Q30
Consensus quality: 157935 bases at least Q20
Insert size: 116494; agarose-fp
Insert size: 165478; sum-of-contigs
Quality coverage: 9.4x in Q20 bases; agarose-fp
Quality coverage: 6.6x in Q20 bases; sum-of-contigs.
NOTE: This is a 'working draft' sequence. It currently
* consists of 15 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 2175: contig of 2175 bp in length
* 1 2176: gap of unknown length
* 2176 2275: gap of unknown length
* 2276 4472: contig of 2197 bp in length
* 4473 4572: gap of unknown length
* 4573 6489: contig of 1917 bp in length
* 6490 6590: gap of unknown length
* 6590 8363: contig of 1774 bp in length
* 8364 8464: gap of unknown length
* 8464 10644: contig of 2181 bp in length
* 10645 10744: gap of unknown length
* 10745 13373: contig of 2629 bp in length
* 13374 13473: gap of unknown length
* 13474 17686: contig of 4213 bp in length
* 17687 17786: gap of unknown length
* 17787 26030: contig of 8244 bp in length
* 26031 26130: gap of unknown length
* 26131 33532: contig of 7402 bp in length
* 33533 33633: gap of unknown length
* 33634 46267: contig of 12635 bp in length
* 46268 46367: gap of unknown length
* 46368 61796: contig of 15429 bp in length
* 61797 61896: gap of unknown length

FEATURES
Source
61897 78516: contig of 16620 bp in length
* 78517 78616: gap of unknown length
* 78617 97532: contig of 18916 bp in length
* 97533 97632: gap of unknown length
* 97633 132687: contig of 35055 bp in length
* 132688 132787: gap of unknown length
* 132788 166878: contig of 34091 bp in length.
Location/Qualifiers
1. 166878
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="8"
/clone="RP11-67H12"
/clone_lib="RP11 human BAC library 11"
1. 2175
/note="assembly_name:Contig22"
2276. 4472
/note="assembly_name:Contig24"
4573. 6489
/note="assembly_name:Contig25"
6590. 8363
/note="assembly_name:Contig28"
8464. 10644
/note="assembly_name:Contig31"
10745. 13373
/note="assembly_name:Contig33"
13474. 17686
/note="assembly_name:Contig34"
17787. 26030
/note="assembly_name:Contig35"
26131. 33532
/note="assembly_name:Contig36
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vector_side:left"
33633. 46267
/note="assembly_name:Contig37"
46368. 61796
/note="assembly_name:Contig38"
61897. 78516
/note="assembly_name:Contig39"
78617. 97532
/note="assembly_name:Contig40
clone_end:T7
vector_side:right"
97633. 132687
/note="assembly_name:Contig41"
132788. 166878
/note="assembly_name:Contig42"
BASE COUNT 41842 a 40044 c 40620 g 42959 t 1413 others
ORIGIN
Query Match 28.4%; Score 262.4; DB 2; Length 166878;
Best Local Similarity 99.6%; Pred. No. 3.2e-54;
Matches 263; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 621 AAAGGCCAAAGCCACAGCCAGAGCCCAAGAGCCAGATTCAAGAAATGAGAGCCACGCC 680
Db 146754 ACAGGCCAAAGCCACAGCCAGAGCCCAAGAGCCAGATTCAAGAAATGAGAGCCACGCC 146813
QY 681 TTGGGGGGCAGCGGTGCAAGTGGGCTTCCCTGGCTGTGCTGACAGGACAGGGTGCCCC 740
Db 146814 TTGGGGGGCAGCGGTGCAAGTGGGCTTCCCTGGCTGTGCTGACAGGACAGGGTGCCCC 146873
QY 741 TGTCAGCCCCCTCCACCTGTGTCTGAATGCAACAGGGGTGTGCGGGGCAACATGAGAG 800
Db 146874 TGTCAGCCCCCTCCACCTGTGTCTGAATGCAACAGGGGTGTGCGGGGCAACATGAGAG 146933
QY 801 CCCCTCACCCCAACTCTCCACTTTACAGAGAGGCCCACTGAAGAGGCCCACTCGGGGT 860
Db 146934 CCCCTCACCCCAACTCTCCACTTTACAGAGAGGCCCACTGAAGAGGCCCACTCGGGGT 146993
QY 861 CACAATTAAGTTGCTGTGCTGAGGA 884

Db 146994 CACAATAAAGTTGCTGCTCAGGA 147017

RESULT 5
AC090442

LOCUS
AC090442

DEFINITION
Homo sapiens chromosome 8 clone RP11-868P18, WORKING DRAFT

AC090442

AC090442

AC090442.2 GI:14389319

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_CANCELLED.

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_CANCELLED.

Homo sapiens.

Homo sapiens.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 181714)

Reichwald, K., Baumgart, C., Blechschmidt, K., Dette, M., Jahn, N., Lehmann, R., Menzel, U., Polley, A., Schilhabel, M.B., Schudy, A., Siddiqui, R., Taudien, S., Wen, G., Rosenthal, A. and Platzer, M.

Chromosome 8 genomic sequence

Unpublished

2 (bases 1 to 181714)

Genome Sequencing Center Jena.

Direct Submission

Submitted (23-FEB-2001) Genome Analysis, Institute of Molecular Biotechnology, Beutenbergstr. 11, Jena 07745, Germany

On Jun 13, 2001 this sequence version replaced gi:13112136.

----- Genome Center

Center: Institute of Molecular Biotechnology

Center code: IMB

Web site: <http://genome.imb-jena.de/>

Contact: gscj-submit@genome.imb-jena.de

----- Project Information

Center project name: H522

Center clone name: RP11-868P18

----- Summary Statistics

Sequencing vector: M13; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.990329

Consensus quality: 167442 bases at least Q40

Consensus quality: 173537 bases at least Q30

Consensus quality: 179477 bases at least Q20

Quality coverage: 5.19 x in Q20 bases; sum-of-contigs

Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality 10.

Quality levels above 40 are expected to have less than 1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

* NOTE: This is a 'working draft' sequence. It currently consists of 15 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 3436: contig of 3436 bp in length

3437 3536: gap of unknown length

3537 10120: contig of 6584 bp in length

10121 10220: gap of unknown length

10221 12191: contig of 1971 bp in length

12192 12291: gap of unknown length

12292 22694: contig of 10403 bp in length

22695 22794: gap of unknown length

22795 42595: contig of 19801 bp in length

42596 42695: gap of unknown length

42696 43816: contig of 1121 bp in length

43817 43916: gap of unknown length

43917 60529: contig of 16613 bp in length

60530 60629: gap of unknown length

60630 66757: contig of 6128 bp in length

66758 66857: gap of unknown length

66858 71578: contig of 4721 bp in length

71579 71678: gap of unknown length

71679 76238: contig of 4560 bp in length

76239 76338: gap of unknown length

76339 88520: contig of 12182 bp in length

88521 88620: gap of unknown length

88621 105835: contig of 17215 bp in length

105836 105935: gap of unknown length

105936 117374: contig of 11439 bp in length

117375 117474: gap of unknown length

117475 163683: contig of 46209 bp in length

163684 163783: gap of unknown length

163784 181714: contig of 17931 bp in length.

location/Qualifiers

1. 181714

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="8"

/clone="RP11-868P18"

1. 4

/note="assembly_fragment"

/vector_side="left"

163784. 181714

misc_feature

/note="Please note that a sequence read was obtained using vectorprimers. This read represents the right end of the clone. However, within this read, the cloning site, e.g. the exact end of insert could not be identified."

BASE COUNT 48155 a 40820 c 40243 g 51096 t 1400 others

ORIGIN

Query Match 28.4%; Score 262.4; DB 2; Length 181714;

Best Local Similarity 99.6%; Pred. No. 3.2e-54;

Matches 263; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 621 AAGGCCAAGCCACAGCCAGAGCCAGCATTCAGAAATGAGAGCCACGCC 680

Db 35142 ACAGGCCAAGCCACAGCCAGAGCCAGCATTCAGAAATGAGAGCCACGCC 35201

QY 681 TTGGGGGGCAGCGTGCAAGTGGGCTTCCCTGGGCTGTGCTGACAGGCGGTGCC 740

Db 35202 TTGGGGGGCAGCGTGCAAGTGGGCTTCCCTGGGCTGTGCTGACAGGCGGTGCC 35261

QY 741 TGTCCAGCCCTCCACCTGTGTCTGAATGCAACAGGGGTGTGGGGGCAACATGAGAG 800

Db 35262 TGTCCAGCCCTCCACCTGTGTCTGAATGCAACAGGGGTGTGGGGGCAACATGAGAG 35321

QY 801 CCCCTCACCCCACTCTCTCCACTTTCAGAGAGGCCCACTGAAGAGCCCACTCGGGGT 860

Db 35322 CCCCTCACCCCACTCTCTCCACTTTCAGAGAGGCCCACTGAAGAGCCCACTCGGGGT 35381

QY 861 CACAATAAAGTTGCTGCTCAGGA 884

Db 35382 CACAATAAAGTTGCTGCTCAGGA 35405

RESULT 6

AC087819

LOCUS
AC087819

DEFINITION
Homo sapiens chromosome 8 clone RP11-507M15 map 8, WORKING DRAFT

AC087819

AC087819.2 GI:13194968

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.

Homo sapiens.

Homo sapiens.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS

1 (bases 1 to 184050)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 8, clone RP11-507M15
Unpublished
2 (bases 1 to 184050)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Boguslavskiy,L., Boukhgalter,B., Brown,A.,
Camarata,J., Campopiano,A., Choepel,Y., Colangelo,M., Collins,S.,
Collimore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodg,S., Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J.,
Gardyna,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Karatas,A., Lacroque,K., Lamazares,R., Landers,T.,
Lehoczky,J., Levine,R., Liu,G., Maclean,C., Macdonald,P.,
Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K.,
McPheeters,R., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R.,
Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rossetti,M.,
Roy,A., Santos,R., Schauer,S., Schupbach,R., Seaman,S., Severy,P.,
Sougnez,C., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J.,
Travers,M., Travis,N., Triggillo,J., Vassiliev,H., Viel,R., Vo,A.,
Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (28-JAN-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Mar 4, 2001 this sequence version replaced gi:12584317.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

TITLE
JOURNAL

COMMENT

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information -----
Center project name: L11344
Center clone name: 507_M15
----- Summary Statistics -----
Sequencing vector: Plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 175465 bases at least Q40
Consensus quality: 179683 bases at least Q30
Consensus quality: 181216 bases at least Q20
Insert size: 170000; agarose-fp
Insert size: 182150; sum-of-contigs
Quality coverage: 7.1 in Q20 bases; agarose-fp
Quality coverage: 6.7 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 20 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 29338: contig of 29338 bp in length
* 29339 29438: gap of 100 bp
* 29439 29457: contig of 19 bp in length
* 29458 29557: gap of 100 bp
* 29558 30186: contig of 629 bp in length
* 30187 30286: gap of 100 bp
* 30287 31083: contig of 797 bp in length
* 31084 31183: gap of 100 bp
* 31184 31978: contig of 795 bp in length
* 31979 32078: gap of 100 bp
* 32079 32748: contig of 670 bp in length
* 32749 32848: gap of 100 bp

FEATURES
source

* 32849 34030: contig of 1182 bp in length
* 34031 34130: gap of 100 bp
* 34131 35157: contig of 1027 bp in length
* 35158 35257: gap of 100 bp
* 35258 72407: contig of 37150 bp in length
* 72408 72507: gap of 100 bp
* 72508 75697: contig of 3190 bp in length
* 75698 75797: gap of 100 bp
* 75798 78108: contig of 2311 bp in length
* 78109 78208: gap of 100 bp
* 78209 82150: contig of 3942 bp in length
* 82151 82250: gap of 100 bp
* 82251 88573: contig of 6323 bp in length
* 88574 88673: gap of 100 bp
* 88674 93173: contig of 4500 bp in length
* 93174 93273: gap of 100 bp
* 93274 101047: contig of 7774 bp in length
* 101048 101147: gap of 100 bp
* 101148 111284: contig of 10137 bp in length
* 111285 111384: gap of 100 bp
* 11385 130797: contig of 19413 bp in length
* 130798 130897: gap of 100 bp
* 130898 152534: contig of 21637 bp in length
* 152535 152634: gap of 100 bp
* 152635 182885: contig of 30251 bp in length
* 182886 182985: gap of 100 bp
* 182986 184050: contig of 1065 bp in length.

Location/Qualifiers

1. 184050

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="8"

/map="8"

/clone_lib="RP11-507M15"
/clone_lib="RP11-11 Human Male BAC"

1. 29338

/note="assembly_fragment"
clone_end:SP6
vector_side:left"

misc_feature
29439. 29457
/note="assembly_fragment"

misc_feature
29558. 30186
/note="assembly_fragment"

misc_feature
30287. 31083
/note="assembly_fragment"

misc_feature
31184. 31978
/note="assembly_fragment"

misc_feature
32079. 32748
/note="assembly_fragment"

misc_feature
32849. 34030
/note="assembly_fragment"

misc_feature
34131. 35157
/note="assembly_fragment"

misc_feature
35258. 72407
/note="assembly_fragment"

misc_feature
72508. 75697
/note="assembly_fragment"

misc_feature
75798. 78108
/note="assembly_fragment"

misc_feature
78209. 82150
/note="assembly_fragment"

misc_feature
82251. 88573
/note="assembly_fragment"

misc_feature
88674. 93173
/note="assembly_fragment"

misc_feature
93274. 101047
/note="assembly_fragment"

misc_feature
101148. 111284
/note="assembly_fragment"

misc_feature
111385. 130797
/note="assembly_fragment"

misc_feature
130898. 152534
/note="assembly_fragment"

misc_feature
/note="assembly_fragment"

misc_feature	152635.	182885	/note="assembly_fragment"
misc_feature	182986.	184050	/note="assembly_fragment"
			clone_end:T7
			vector_side:right"
BASE COUNT	49622	a 42339	c 42431 g 47728 t 1930
ORIGIN			others

Query Match	28.48;	Score 262.4;	DB 2;	Length 184050;
Best Local Similarity	99.6%;	Pred. No. 3.2e-54;		
Matches 263; Conservative	0;	Mismatches 1;	Indels 0;	Gaps 0;

QY	621	AAAGGCCAAAGCCACACAGCCAGAGGCCAAGAAAGCCAGATTTCAAGAAATGAGGAGCCACGCCC	680
Db	123140	ACAGGCCAAAGCCACACAGCCAGAGGCCAAGAAAGCCAGATTTCAAGAAATGAGGAGCCACGCCC	123199
QY	681	TTGGGGGGCGCAGGTGCAGAAAGTGGGCTTCCCTGGGCTGTGCTGCAGGCGACAGGGTGGCCCC	740
Db	123200	TTGGGGGGCGCAGGTGCAGAAAGTGGGCTTCCCTGGGCTGTGCTGCAGGCGACAGGGTGGCCCC	123259
QY	741	TGTCACAGCCCTCCACCTGTGTCTGAATGCAACAGAGGGTGTGCGGGGGCAACATGAGAG	800
Db	123260	TGTCACAGCCCTCCACCTGTGTCTGAATGCAACAGAGGGTGTGCGGGGGCAACATGAGAG	123319
QY	801	CCCCCTACCCCCCAACTCTCCACTTTTCAGGAGGCCCCCAGTGAAGAGCCCCCACCCTGGGGGT	860
Db	123320	CCCCCTACCCCCCAACTCTCCACTTTTCAGGAGGCCCCCAGTGAAGAGCCCCCACCCTGGGGGT	123379
QY	861	CACAATTAAGTTGCCTGGTCAAGGA	884
Db	123380	CACAATTAAGTTGCCTGGTCAAGGA	123403

RESULT 7	LOCUS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE	ORGANISM	REFERENCE	AUTHORS
AC090471/c	AC090471	Homo sapiens chromosome 8 clone RPL1-868P18 map 8, WORKING DRAFT SEQUENCE, 10 unordered pieces.	AC090471	AC090471.6	GI:14249070	HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLLTOP.	Homo sapiens.	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.	Birren,B., Linton,L., Nusbaum,C. and Lander,E.
								1 (bases 1 to 184444)	
								Homo sapiens chromosome 8, clone RPL1-868P18	
								unpublished	
								2 (bases 1 to 184444)	
								Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.	

REFERENCE	AUTHORS	TITLE	JOURNAL	REFERENCE
1 (bases 1 to 184444)	Birren, B., Linton, L., Nusbaum, C. and Lander, E.	Homo sapiens chromosome 8, clone RP11-868P18	Unpublished	
2 (bases 1 to 184444)	Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Bouckgalter, B., Brown, A., Camarata, J., Campopiano, A., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferrelta, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Hearford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Karatas, A., Larocque, K., Lamazares, R., Landers, T., Lehoczký, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Mcpheeters, R., Meldrim, J., Meneus, L., Mlnova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Plerre, N., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupback, R., Seaman, S., Severy, P., Sougnuez, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.			
Direct Submission				

JOURNAL
COMMENT

Submitted (23-FEB-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 30, 2001 this sequence version replaced gl:14196391.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information -----

Center project name: L11354
Center clone name: 868_P_18

Summary Statistics

Chemistry: Dye-terminator Big Dye 3, 100% of reads
sequencing Vector: Plasmid; n/a; 100% of reads

```
Assembly program: Phrap; version 0.960731
```

consensus quality:	180698	bases at least Q40
consensus quality:	182469 <td>bases at least Q30</td>	bases at least Q30

Consensus quality: 183168 bases at least Q20

```
Insert size: 183000; agarose-tp
Insert size: 183544; sum-of-contigs
```

Quality coverage:	15.1	in Q20 bases; agarose-fp
Quality coverage:	15.1	in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 10 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

*	1	24646:	contlg of 24646 bp	in length
*	24647	24746:	gap of	100 bp
*	24747	28060:	contlg of 3314 bp	in length
*	28061	28160:	gap of	100 bp
*	28161	33796:	contlg of 5636 bp	in length
*	33797	33896:	gap of	100 bp
*	33897	41745:	contlg of 7849 bp	in length
*	41746	41845:	gap of	100 bp
*	41846	52788:	contlg of 10943 bp	in length
*	52789	52888:	gap of	100 bp
*	52889	68506:	contlg of 15618 bp	in length
*	68507	68606:	gap of	100 bp
*	68607	105368:	contlg of 36762 bp	in length
*	105369	105468:	gap of	100 bp
*	105469	133826:	contlg of 28358 bp	in length
*	133827	133926:	gap of	100 bp
*	133927	170402:	contlg of 36476 bp	in length
*	170403	170502:	gap of	100 bp
*	170503	184444:	contlg of 13942 bp	in length

FEATURES
source

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/organism="Homo sapiens"  
/db_xref="taxon:9606"  
/chromosome="8"  
/map="8"  
/clone="RP11-868p18"  
/clone_lib="RPCI-11 Human Male BAC  
l. 24646  
/note="assembly_fragment  
clone_end:Sp6  
vector_side:left"  
24747..28060  
/note="assembly_fragment"  
28161..33796  
/note="assembly_fragment"  
33897..41745  
/note="assembly_fragment"  
41846..52788  
/note="assembly_fragment"
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misc_feature      52889..68506
                  /note="assembly_fragment"
misc_feature      68607..105368
                  /note="assembly_fragment"
misc_feature      105469..133826
                  /note="assembly_fragment"
misc_feature      133927..170402
                  /note="assembly_fragment"
misc_feature      170503..184444
                  /note="assembly_fragment"
                  clone_end:T7
                  vector_side:right"

BASE COUNT      52113 a 41006 c 41428 g 48985 t 912 others
ORIGIN

Query Match      28.4%; Score 262.4; DB 2; Length 184444;
Best Local Similarity 99.6%; Pred. No. 3.2e-54;
Matches 263; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 621 AAAGGCCAAGCCACAGCCAGAGCCAGATTCAGAAATGAGAGCCACGCC 680
    |
Db 10772 ACAGGCCAAGCCACAGCCAGAGCCAGATTCAGAAATGAGAGCCACGCC 10713
    |

QY 681 TTGGGGGGCAGCGGTGCAAAAGTGGGCTTCCCTGGGCTGTGCTGCAGCAGCAGGCTGCCCC 740
    |
Db 10712 TTGGGGGGCAGCGGTGCAAAAGTGGGCTTCCCTGGGCTGTGCTGCAGCAGCAGGCTGCCCC 10653
    |

QY 741 TGTCCAGCCCCCTCCACCTGTGTCTGATGCAACAGGGGCTTGGCGGGCAACATGAGAG 800
    |
Db 10652 TGTCCAGCCCCCTCCACCTGTGTCTGATGCAACAGGGGCTTGGCGGGCAACATGAGAG 10593
    |

QY 801 CCCCTCACCCCCCACTCTCCACTTTTGAAGAGGGCCCCCAGTGAAGAGCCCCCAGCTCGGGGT 860
    |
Db 10592 CCCCTCACCCCCCACTCTCCACTTTTGAAGAGGGCCCCCAGTGAAGAGCCCCCAGCTCGGGGT 10533
    |

QY 861 CACAATAAAGTTGCTGTGTCAGGA 884
    |
Db 10532 CACAATAAAGTTGCTGTGTCAGGA 10509
    |

RESULT 8
HS833B7/c      86574 bp DNA linear PRI 12-DEC-1999
LOCUS
DEFINITION
    Human DNA sequence from clone CTA-833B7 on chromosome 22q12.3-13.2
    Contains the NCF4 gene for cytosolic neutrophil factor 4 (40kd),
    the 5' part of the CSF2RB gene for granulocyte-macrophage
    low-affinity colony stimulating factor 2 receptor beta, ESTs, STSS
    and GSSs, complete sequence.
ACCESSION
    AL008637
    AL008637.1 GI:3136000
VERSION
    HTG; colony stimulating factor; CSF2RB; cytosolic neutrophil
    factor; NCF4.
KEYWORDS
    Homo sapiens.
SOURCE
    Homo sapiens.
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
    1 (bases 1 to 86574)
    Burton, J.
    Direct Submission
    Submitted (08-DEC-1999) Sanger Centre, Hinxton, Cambridgeshire,
    CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
    requests: clonerequests@sanger.ac.uk
    On May 18, 1998 this sequence version replaced gi:2578146.
    This sequence has been finished according to sequence map criteria
    as follows. An attempt is made to resolve all sequencing problems,
    such as compressions and repeats, but not necessarily within known
    annotated human repeat sequence elements (e.g. Alu). Where the
    sequence is ambiguous, there is an annotation using the 'unsure'
    feature key.
    This sequence was generated from part of bacterial clone contigs of
    human chromosome 22, constructed by the Sanger Centre Chromosome 22
    Mapping Group. Further information can be found at
    http://www.sanger.ac.uk/HGP/Chr22

```

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep CTA-833B7 is from the human BAC library described in U-J. Kim et al. (1996) Genomics 34, 213-218.

VECTOR: pBelBAC11

IMPORTANT: This sequence is not the entire insert of clone CTA-833B7 it may be shorter because we only sequence overlapping sections once, or longer because we arrange for a small overlap between neighbouring submissions.

The true right end of clone CTF22-24E5 is at 100 in this sequence. The start of this sequence overlaps with sequence Z82185.

FEATURES

source

Location/Qualifiers

1..86574

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="22"

/map="q12.3-13.2"

/clone="CTA-833B7"

/clone_lib="CIT978SK-A2"

1..176

/note="AluSp repeat: matches 138..313 of consensus"

862..903

/note="MIR repeat: matches 212..254 of consensus"

1791..2095

/note="Aluub repeat: matches 1..307 of consensus"

complement(2600..3449)

/note="match: GSS: Em:AQ749444"

3359..3522

/note="MER58A repeat: matches 43..207 of consensus"

3532..3838

/note="AluSx repeat: matches 1..308 of consensus"

4030..4288

/note="MLT1J repeat: matches 1..261 of consensus"

complement(4402..4679)

/note="match: GSS: Em:AQ321484"

5050..5195

/note="LTR41 repeat: matches 4..144 of consensus"

5291..5370

/note="L1ME1 repeat: matches 5526..5607 of consensus"

5371..5676

/note="AluSx repeat: matches 1..310 of consensus"

5677..6252

/note="L1ME1 repeat: matches 5607..6163 of consensus"

6302..6485

/note="L2 repeat: matches 2078..2265 of consensus"

6873..7196

/note="54 copies 6 mer gtgtgt 66 conserved"

6874..7197

/note="27 copies 12 mer 66 conserved"

6879..7190

/note="13 copies 24 mer 67 conserved"

6884..7195

/note="156 copies 2 mer tg 67 conserved"

6886..7193

/note="22 copies 14 mer 67 conserved"

6907..7194

/note="9 copies 32 mer 69 conserved"

7203..7332

/note="L2 repeat: matches 2602..2750 of consensus"

7347..7532

/note="MIR repeat: matches 2..191 of consensus"

7537..7640

/note="MIR repeat: matches 158..262 of consensus"

8152..8228

TITLE Haugen, E.D.
Direct Submission
JOURNAL Submitted (01-SEP-2001) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 149940)
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Submitted (24-MAR-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
COMMENT On Mar 24, 2002 this sequence version replaced gl.18087686.

----- Genome Center

Center: University of Washington Genome Center
Center Code: UWGC

Web site: <http://www.genome.washington.edu>
Contact: uwgchgs@u.washington.edu

Drafting Center: BCM

----- Project Information

Center project name: chr-3
Center clone name: RP11-56P22 (bc0176)

----- Summary Statistics

Sequencing vector: M13; 108821: 52% of reads
Sequencing vector: plasmid; 48% of reads
Sequencing vector: plasmid; 108752: 0% of reads
Chemistry: Dye-terminator ET; 17% of reads
Chemistry: Dye-terminator Big Dye; 82% of reads
Chemistry: Dye-terminator Big Dye; 82% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 149854 bases at least Q40
Consensus quality: 149935 bases at least Q30
Consensus quality: 149940 bases at least Q20
Insert size: 149940; sum-of-contigs
Quality coverage: 8.3x in Q20 bases; sum-of-contigs

----- Overlapping Sequences:

5': RP11-237P23 (UWGC:bc0783) AC112223
3': RP11-114C6 (UWGC:bc0623) AC108486

----- Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program. All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than 1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., Phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

----- Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both insert and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400-800 bp) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

ECORI

HindIII

BglII

SeqDerMap FngPrnt SeqDerMap FngPrnt SeqDerMap FngPrnt

8696	8608	3232	3238	10777	10725
6	<800	6382	6740	2067	2070
1938	1915	512	<800	8879	9137
2860	2886	449	<800	4717	4762
665	<800	3551	3535	1161	1132
1137	1166	66	<800	3316	3323
3903	3847	4029	4016	7117	7092
357	<800	1886	1892	601	<800
2519	2497	379	<800	14637	14477
212	<800	5099	5054	23604	23653
3478	3509	2949	3028	2916	2926
2469	2497	5004	5054	2176	2167
873	900	1085	1067	556	<800
279	<800	1671	1632	3689	3724
5376	5312	393	<800	2880	2926
2045	2067	8291	8472	7884	7435
1963	1915	1962	1892	565	<800
1172	1166	7255	7042	3478	3460
1272	1307	2118	2170	2658	2665
1083	1037	1169	1168	741	744
9619	9736	1330	1292	2895	2926
1642	1625	235	<800	11488	11677
199	<800	1294	1292	1498	1521
3489	3509	857	901	2400	2416
4629	4644	2838	2805	5145	5305
8423	8608	4757	4713	1202	1162
3204	3186	2331	2337	30	<800
6821	6882	1896	1892	1839	1807
1033	1037	761	768	1978	1930
4701	4644	1612	1632	2469	2416
17125	16431	2232	2170	2637	2665
2527	2497	6677	6740	808	830
2974	2886	201	<800	6888	6667
3166	3186	677	<800	4739	4762
4025	4005	265	<800	1346	1323
14230	14884	3833	3824	2429	2416
5750	5785	7946	7558	12	<800

-----	-----	-----	-----
279	<800	2301	2337
-----	-----	-----	-----
1323	1307	2753	2805
-----	-----	-----	-----
12	<800	490	<800
-----	-----	-----	-----
2101	2067	177	<800
-----	-----	-----	-----
4290	4236	6836	6740
-----	-----	-----	-----
1154	1166	6767	6740
-----	-----	-----	-----
766	801	5595	5684
-----	-----	-----	-----
6886	6882	5959	5945
-----	-----	-----	-----
7307	7318	10600	10609
-----	-----	-----	-----
		5616	5684
		5062	5054
		111	<800
		4421	4433
		1890	1892
		4176	4156

FEATURES
source

Location/Qualifiers
1. 149940
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP11-56P22"
/clone_lib="RP11 human BAC library 11"

Query Match 13.08; Score 120.4; DB 9; Length 149940;

Best Local Similarity 61.58; Pred. No. 5.1e-19;

Matches 368; Conservative 0; Mismatches 156; Indels 74; Gaps 8;

```
QY 317 TTCTCTCCCACTTACTTTCCAGCTCCGGCTGGCTCAGAGCCGCTGTTCTCAGTGCC 376
      ||||| ||| ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 79005 TTCTCTCCCACTTACTTTTAACTCTGAGCT-GCTCCAAATCCCTGTTCTTCACTGACT 79063

QY 377 AGGAACGTTATGAGCATCAGACCTAACCTGGAGAGGAGGAGGAGGAGGAGGAGGAGC 436
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 79064 AAAGATGTTATGAACCTTTAGCCCTAACCTGGAGAGGAGGAGGAGGAGGAGGAGGAG 79123

QY 437 AGGAGGAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGC 496
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 79124 ACGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 79159

QY 497 AAAGCCCTGTCAACAGTCAAAAGGCTGGTCCCAAGAGAGGAGGAGGAGGAGGAGGAG 556
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 79160 AAACCTCTATCAACATGTCAAAAGGTGGGGGAGCCCAAGAGGAGGAGGAGGAGGAG 79218

QY 557 AAAAAAGCTGGAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGC 616
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 79219 GCGAGGAGCTGGAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 79268

QY 617 TGAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 676
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 79269 TGAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 79328

QY 677 -----CGCCTTGGGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 706
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 79329 GGAAGTGCCTGACTGGGCTGATCAAGAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 79388

QY 707 TTCCCTGGGCTGTGCTGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 763
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
```

```
Db 79389 TTCCCTGGGCTGTGCTGATCATGGGGTATACCCCTACCTTGCCCTTCCATCTGAGTC 79448
QY 764 TGAATGCAACAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 819
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 79449 TAAATAGAGTGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 79508
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RESULT 11

XLNUPLR

LOCUS XLNUPLR 646 bp mRNA linear VRT 10-FEB-1999

DEFINITION

ACCESION Y00204.1 GI:64938

VERSION Y00204.1

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

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FEATURES
source

Location/Qualifiers
1. 646
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polyA_signal 222 a 121 c 174 g 129 t
BASE COUNT 222 a 121 c 174 g 129 t
ORIGIN

Query Match

Best Local Similarity

Matches 224; Conservative

10.88; Score 99.6; DB 5; Length 646;
57.48; Pred. No. 4.8e-14;
0; Mismatches 154; Indels 12; Gaps 2;

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QY 69 GACCTGCTCTGCGGCTGCGAGCTCAGTCAGAGAGGAGGAGGAGGAGGAGGAGGAGGAG 128
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 29 GTCCCTTATTTGGGGGTGTAACATGAGCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 88

QY 129 GCTGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 185
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Db 89 AGACGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 148

QY 186 AGCCAAAGAGGAGATGCATCGGCTGAGATCCTGCCCCAGCAAAACAGAGAGGAGCAAGAA 245
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Db 149 GGCAAGAGATGAGTTCACATAGTAGAAT-----AGTTACACAGAGAGGAGGAGGAAA 199
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QY 724 CAGGCACAGGCTGCCCTGTCACGCCCTCCACCTG 759
::: : :::: : ||||| |||||
Db 1081 RRRRRRRRRRRRRATCGCAGCTCCTCGACCTG 1046

RESULT 14
AC125180

LOCUS AC125180 198421 bp DNA linear HTG 20-JUN-2002
DEFINITION Mus musculus chromosome UNK clone RP23-322K1, WORKING DRAFT
SEQUENCE, 30 unordered pieces.

AC125180
AC125180.1 GI:21490699
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 198421)
McPherson,J.D. and Waterston,R.H.
The sequence of Mus musculus clone
Unpublished
2 (bases 1 to 198421)
McPherson,J.D. and Waterston,R.H.
Direct Submission
Submitted (20-JUN-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA

REFERENCE
AUTHORS
TITTLE
JOURNAL
REFERENCE
AUTHORS
TITTLE
JOURNAL
COMMENT

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Contact: submissions@watson.wustl.edu
----- Project Information -----
Center project name: M_BA0322K01
----- Summary Statistics -----
Sequencing vector: M13; 0%
Sequencing vector: plasmid: 100%
Chemistry: Dye-primer ET; 0% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 186437 bases at least Q40
Consensus quality: 189477 bases at least Q30
Consensus quality: 191697 bases at least Q20
Insert size: 203000; agarose-fp
Insert size: 196692; sum-of-contigs
Quality coverage: 4.53 in Q20 bases; agarose-fp
Quality coverage: 4.25 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 30 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1076: contig of 1076 bp in length
* 1077 1176: gap of unknown length
* 1177 2506: contig of 1330 bp in length
* 2507 2606: gap of unknown length
* 2607 4336: contig of 1730 bp in length
* 4337 4436: gap of unknown length
* 4437 5587: contig of 1151 bp in length
* 5588 5687: gap of unknown length
* 5688 6909: contig of 1222 bp in length
* 6910 7009: gap of unknown length
* 7010 8416: contig of 1407 bp in length
* 8417 8516: gap of unknown length
* 8517 10353: contig of 1837 bp in length
* 10354 10453: gap of unknown length
* 10454 12880: contig of 2427 bp in length
* 12881 12980: gap of unknown length

* 12981 15704: contig of 2724 bp in length
* 15705 15804: gap of unknown length
* 15805 18535: contig of 2731 bp in length
* 18536 18635: gap of unknown length
* 18636 21653: contig of 3018 bp in length
* 21654 21753: gap of unknown length
* 21754 23575: contig of 1822 bp in length
* 23576 23675: gap of unknown length
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* 26532 26631: gap of unknown length
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* 30436 30535: gap of unknown length
* 30536 35803: contig of 5268 bp in length
* 35804 35903: gap of unknown length
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* 41091 41190: gap of unknown length
* 41191 46181: contig of 4991 bp in length
* 46182 46281: gap of unknown length
* 46282 53388: contig of 7107 bp in length
* 53389 53488: gap of unknown length
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* 73617 80218: contig of 6602 bp in length
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* 80319 87536: contig of 7218 bp in length
* 87537 87636: gap of unknown length
* 87637 97222: contig of 9586 bp in length
* 97223 97322: gap of unknown length
* 97323 105689: contig of 8367 bp in length
* 105690 105789: gap of unknown length
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* 123928 138453: contig of 14526 bp in length
* 138454 138553: gap of unknown length
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* 160962 161061: gap of unknown length
* 161062 197994: contig of 36933 bp in length
* 197995 198094: gap of unknown length
* 198095 198421: contig of 327 bp in length.

FEATURES
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location/Qualifiers
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/note="assembly_name:Contig23"
5670. .8287
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14273. .20234
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20335. .28899
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61366. .68507
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68608. .84968
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Query Match	9.0%;	Score 83.2;	DB 2;	Length 190604;
Best Local Similarity	61.6%;	Pred. No. 8.6e-10;		
Matches 133; Conservative	0;	Mismatches 83;	Indels 0;	Gaps 0;

[illegible]

Search completed: January 24, 2003, 18:22:30
Job time : 46614 secs

GenCore version 5.1.3
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 23, 2003, 14:00:24 ; Search time 310 Seconds
(without alignments)
6712.408 Million cell updates/sec

Title: US-09-844-864-16
Perfect score: 924
Sequence: 1 cagccgcgtctctgcccgg.....ttcgccgccaagcttatg 924

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues
Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_101002:*

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2: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1981.DAT:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	858	92.9	1048	21	AAA72432	Human nucleic acid
2	342.8	37.1	1019	21	AAD00296	Mouse oocyte-spect
3	337	36.5	423	23	AAD23440	Human lung tumour-
4	171.6	18.6	86574	24	ABK83560	Human CDNA differe
5	131.6	14.2	1481	21	AAD00297	Mouse ovary-specif
6	72.4	7.8	416	22	AA101804	Human reproductive
7	72.4	7.8	416	23	ABL97097	Human testicular a
8	70.2	7.6	49999	20	AAZ23891	Murine LOBO genomi
9	70.2	7.6	49999	20	AAZ23896	Murine LOBO homolo

10	69.2	7.5	33923	22	AAK67071	Human immune/haema
11	68.8	7.4	341	23	AAS90676	DNA encoding novel
12	67.8	7.3	512	24	ABQ54697	Human ovarian anti
13	66.4	7.2	291	23	AAS68926	DNA encoding novel
14	65.8	7.1	1680	23	AAS68538	DNA encoding novel
15	65.2	7.1	8165	22	AAK82672	Human immune/haema
16	64	6.9	510	23	AAS69539	DNA encoding novel
17	64	6.9	510	23	AAS71141	DNA encoding novel
18	64	6.9	510	23	AAS90687	DNA encoding novel
19	63.8	6.9	591	23	AAS70521	DNA encoding novel
20	63.8	6.9	591	23	AAS70706	DNA encoding novel
21	63.8	6.9	591	23	AAS90721	DNA encoding novel
22	63.8	6.9	1416	23	AAS67163	DNA encoding novel
23	63.4	6.9	575	22	ABA50472	DNA encoding novel
24	63.4	6.9	575	22	ABA68422	Human breast cell
25	63.4	6.9	575	22	ABA35413	Human foetal liver
26	63.4	6.9	575	22	AAK16793	Human brain expres
27	63.4	6.9	575	22	AAK42567	Human bone marrow
28	63.4	6.9	575	22	AAI23315	Probe #13248 for g
29	63.4	6.9	575	22	AAI48636	Probe #17322 used
30	63.4	6.9	575	22	AAI08956	Probe #8947 used t
31	63.4	6.9	575	24	ABS16615	Human genome-derl
32	63.4	6.9	1969	22	ABA45341	Human breast cell
33	63.4	6.9	1969	22	ABA55830	Human foetal liver
34	63.4	6.9	1969	22	ABA25506	Probe #3972 for ge
35	63.4	6.9	1969	22	AAK04048	Human brain expres
36	63.4	6.9	1969	22	AAK29533	Human bone marrow
37	63.4	6.9	1969	22	AAI14105	Probe #4038 for ge
38	63.4	6.9	1969	22	AAI35486	Probe #4172 used t
39	63.4	6.9	1969	22	AAI03958	Probe #3949 used t
40	63.4	6.9	1969	24	ABS04084	Human genome-derl
41	63.4	6.9	51259	18	AAK83007	Human immune syste
42	63.2	6.8	14798	24	ABJ33033	Partial mouse WRN
43	63	6.8	1080	24	ABJ99537	Human immune syste
44	62.6	6.8	654	23	AAS68925	Mouse ischaemic co
45	62.4	6.8	575	22	ABA63148	DNA encoding novel

ALIGNMENTS

RESULT 1	
AAA72432	
ID AAA72432 standard; CDNA; 1048 BP.	
AC AAA72432;	
DT 19-DEC-2000 (first entry)	
DE Human nucleic acid-binding protein NuABP-51 cDNA.	
DE Human nucleic acid-binding protein; NuABP; agonist; antagonist; EST;	
KW expressed sequence tag; drug screening; recombinant expression; antibody;	
KW reproductive disorder; infertility; immunological disorder;	
KW neurological disorder; cell proliferative disorder; cancer; tumour; ss.	
OS Homo sapiens.	
PN WO200044900-A2.	
PD 03-AUG-2000.	
PF 28-JAN-2000; 2000WO-US02237.	
PR 29-JAN-1999; 99US-0117904.	
PR 29-JAN-1999; 99US-0117905.	
PA (INCY-) INCYTE PHARM INC.	
PI Tang YT, Lal P, Hillman JL, Yue H, Azimzai Y, Lu AMD, Baughn MR;	
PI Tran B, Shih LL, Au-Young JL;	
PI WPI; 2000-499332/44.	

CC cDNA clone O1-236. It is derived from mouse 2-cell embryo cDNA library
CC and expressed in the oocytes of intermediate size type 3a follicles and
CC all type 3b follicles. This clone is used to screen and identify the
CC mouse Npm2 gene. It is homologous to Xenopus laevis nucleoplasm (Xnm2)
CC expressed exclusively in eggs. It provides in vitro and in vivo reagents
CC for studying ovarian development and function. This sequence has
CC gynaecological and contraceptive activity. Agents which modulate O1-180,
CC O1-184 and O1-236 may be used to treat cell proliferative or degenerative
CC disorders, associated with abnormal expression of these ovary specific
CC genes. This ovary-specific sequence can be used as reagents to evaluate
CC potential contraceptives, to block ovulation in a reversible manner.
CC It is also used to screen for genetic mutations in signalling pathways,
CC that are associated with some forms of human infertility or
CC gynaecological cancers.

XX
SQ Sequence 1019 BP; 334 A; 243 C; 265 G; 177 T; 0 other;

Query Match 37.1%; Score 342.8; DB 21; Length 1019;
Best Local Similarity 66.4%; Pred. NO. 1.5e-72;
Matches 603; Conservative 0; Mismatches 262; Indels 43; Gaps 6;

OY 1 CAGCCCGCTTCTGCGCGGACCATGATCTCAGTAGCCGACAGTACAGCGAGAAAG 60
Db 132 CACCAAGCCGCCCTGTAATCGACATGAGTCGCCACAGCACAGCAGCGTGACCGAAACC 191
OY 61 GCAGTGACGACCGTCTGCGGGCTGCGAGCTCAGTCAGAGAGAGCGGACCTTTC 120
Db 192 ACAGCAAAAAACATGCTCGGGGTAGTGAACCTCATTCAGGAAAGCAGACTTGCACCTTT 251
OY 121 AGACCCACAGCTGGAGGGGAGCAGAGCTGACGCTGTTGCTTACATGATTGCTGGGG 180
Db 252 AGAGGCCAAGCGGAGAGAGAGCAGCAGCTGTAACCTTCTGCTCAGCAGCAGTCTGGGG 311
OY 181 GAGAAAGCCAAAGAGAGATGATCGCTGAGATCTGCCCCCAGCAAAACAGAGAGAC 240
Db 312 GAGCAAGCCAAAGAGAGAGTGAACCGTGTGAGTCTCTCC-----AGGAA 359
OY 241 AAGAAGATGACCGCGTCACCATTTGCTCAGTCAGGCTCAGTCTCCCATGCTCTCC 300
Db 360 GCGAGAAACCACCAATCACTATTGCTACGCTGAAGCATCAGTCTGCCCATGCTCACT 419
OY 301 ATGCTAGAGTGCAGCTTCTCCCGCAGTTACTTCCAGCTCCGGCTGGCTCAGAGACC 360
Db 420 GTGTCAAGTATAGACTTCTCTCCAGTAACCTTTCCGCTCAGACCTGGCTCAGACCT 479
OY 361 GTGTCTCAGTGGCAGGAACCTTATGAAGATCAGACCTAACCTGGAGAGAGAGAG 420
Db 480 GTGTCTCAGTGGCTGGAATGTTATGAGACTTCCGACCTGACCTGGAGATGACGAG 539
OY 421 GAAGAAGAAGGAGAGAGAGAGAGAGAGAGATGATGAGATGAGATGACAGAT 480
Db 540 GAA-----GAGGAGGAAGAGAGAGAGAGAGAGATGAGATGAGATGACAGAT 587
OY 481 ATATCTGAGAGAGCAAAAGCCCTGTCAAAAGTCAAAAGGCTGGTGGCCCGAGAGCAG 540
Db 588 ATATCGCTAGAGAGAG---ATACCTGTCAAAAGTCAAAAGGCTGGTGGCCCGAGAGCAG 644
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Db 645 ATGAGCATAGCAAGAAAAAAGAGGTGAAAAAAGAGAGATGAACAGTAGTAGGCCCC 704
OY 595 AGCGTTAGAGACAGAGCCCTGTGAAAAAGGCCAAAGCCAGAGCCAGAGAGAGAGAG 654
Db 705 AGCCCTCAGAGACAGAGTCCCTGGAAGAGAGAGAAATCTACACCCAGAGAGAGAGAG 764
OY 655 GCATTCAAGAAATGAG 714
Db 765 GTGACCAAGAAATGA--CCTCATCTTACCATCTTCTGCTCAAGGAGAGATGTCCAGCA 822
OY 715 GCTGTGCTGACAG 774
Db 823 GCTGTGCTGAG 874

OY 775 GGGGTGTGCGGGGCAACATGAGAGCCCTTCACCCCAACTCTTCACCTTTCAGAGGCC 834
Db 875 AGGTGTGTGCTGTAAACCTGTAAACCCAGCCCTTCAGTTCCGAGAGTTTGTGAAG 934
OY 835 CCCAGTGAAGAGCCCCACCTCGGGGTGACATTAAGTTGCTGTCAGAGAAAAA 894
Db 935 AGCCCCCAGCAAGTTCGCTAGGGCCACAAATAAATTGCATGATCAGAAAAA 994
OY 895 AAAAAAA 902
Db 995 AAAAAAA 1002

RESULT 3
AAD23440
ID AAD23440 standard; cDNA; 423 BP.

XX
AC AAD23440;
XX
DT 26-FEB-2002 (first entry)

XX Human lung tumour-specific 54589.1 cDNA.

XX Human; lung tumour protein; immunostimulant; cytostatic; gene therapy;
KM antisense-therapy; vaccine; immune response; lung cancer; 54589.1; ss.

OS Homo sapiens.

PN WO200172295-A2.

PD 04-OCT-2001.

PF 28-MAR-2001; 2001WO-US09991.

PR 29-MAR-2000; 2000US-0538037.

PR 05-JUN-2000; 2000US-0588937.

PR 18-AUG-2000; 2000US-0640878.

PR 22-SEP-2000; 2000US-234517P.

PR 01-NOV-2000; 2000US-0704512.

PR 14-DEC-2000; 2000US-0738973.

XX (CORI-) CORIXA CORP.

XX Reed SG, Lodes MJ, Mohamath R, Secrist H, Benson DR, Indirias CY;

PI Henderson RA, Fling SP, Algate PA, Elliot M, Munton J, Kalos MD;

XX WPI; 2001-639201/73.

PT New human lung-specific polynucleotides and polypeptides for the
PT diagnosis and treatment of disease e.g. lung cancer -

XX Claim 1; Page 302; 378pp; English.

XX The invention relates to isolated lung tumour-specific proteins and
CC their corresponding cDNA molecules. Lung tumour-specific proteins and
CC their antigen-presenting cells are useful for stimulating and/or
CC expanding T cells specific for a tumour protein, and for inhibiting
CC the development of cancer. The invention also relates to a composition
CC useful for stimulating an immune response, and for treating cancer. The
CC lung tumour specific oligonucleotide is useful in gene therapy and for
CC diagnosis, detection and treatment of lung cancer. The present sequence
CC is a cDNA encoding human lung tumour-specific protein.

XX
SQ Sequence 423 BP; 84 A; 138 C; 125 G; 76 T; 0 other;

Query Match 36.5%; Score 337; DB 23; Length 423;
Best Local Similarity 100.0%; Pred. NO. 2.7e-71;
Matches 337; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CAGCCCGCTTCTGCGCGGAGCCATGATCTCAGTAGCGCAGTAGAGAGAGAGAG 60
Db 87 CAGCCCGCTTCTGCGCGGAGCCATGATCTCAGTAGCGCAGTAGAGAGAGAGAG 146

QY	61	GCAGTGCAGCACCCTGCTCTGTGGGGCTGCGAGCTCAGTCAAGAGAGGGCGGACTTGGACCTTC	120
Db	147	GCAGTGCAGCACCCTGCTCTGTGGGGCTGCGAGCTCAGTCAAGAGAGGGCGGACTTGGACCTTC	206
QY	121	AGACCCCAAGCTGAGGGGGAAGCAGAGCTGCAGGCTGTGCTTCATACGATTGTGCTTGGGG	180
Db	207	AGACCCCAAGCTGAGGGGGAAGCAGAGCTGCAGGCTGTGCTTCATACGATTGTGCTTGGGG	266
QY	181	GAGAAAGCCAAAGAGAGAGATGCATCGCGTGGAGATCCTGCCCCAGCAAAACAGAGAGAC	240
Db	267	GAGAAAGCCAAAGAGAGAGATGCATCGCGTGGAGATCCTGCCCCAGCAAAACAGAGAGAC	326
QY	241	AAGAAGATGCAGCCGGTCAACCATTGCTCACTCCAGGCTCAAGTCTCCCATGGTCTCC	300
Db	327	AAGAAGATGCAGCCGGTCAACCATTGCTCACTCCAGGCTCAAGTCTCCCATGGTCTCC	386
QY	301	ATGGTAGAGTGCAGCTTCTCCCCAGTTACTTTCC	337
Db	387	ATGGTAGAGTGCAGCTTCTCCCCAGTTACTTTCC	423

RESULT 4
ABK83560/c
ID ABK83560 standard; cDNA; 86574 BP.

AC ABK83560;

DT 14-AUG-2002 (first entry)

DE Human cDNA differentially expressed in granulocytic cells #131

AA Human; ss; granulocytic cell; DNA chip; bacterial infection;
 KW viral infection; parasitic infection; protozoal infection;
 KW fungal infection; sterile inflammatory disease; psoriasis;
 KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;
 KW cardiac reperfusion injury; renal reperfusion injury; ARDS;
 KW adult respiratory distress syndrome; inflammatory bowel disease;
 KW Crohn's disease; ulcerative colitis; periodontal disease;
 KW granulocyte activation; chronic inflammation; allergy.

OS Homo sapiens.

PN W0200228999-A2.

PD 11-APR-2002.

PF 03-OCT-2001; 2001WO-US30821.

PR 03-OCT-2000; 2000US-237189P.

PA (GENE-) GENE LOGIC INC.

PI Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;

DR WPI; 2002-435328/46.

PT Detecting granulocyte activation by detecting differential expression
PT of genes associated with granulocyte activation, which serves as
PT diagnostic markers that is useful for monitoring disease states and
PT drug toxicity -

PS Claim 1; SEQ ID No 131; 114pp; English.

The invention relates to detecting (M1) granulocyte (GC) activation (GCA), by detecting the level of expression of gene(s) (Gs) identified by DNA chip analysis as given in the specification, and comparing the expression level to an expression level in an unactivated GC, where differential expression of Gs is indicative of GCA. Also included are modulating (M2) GA by contacting GC with an agent that alters the expression of at least one gene in Gs; (2) screening (M3) for an agent capable of modulating GCA or an inflammation (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease using the

gene expression profile; (3) detecting (M4) an inflammation (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease, by detecting the level of expression in a sample of the tissue of gene(s) from Gs, where the level of expression of the gene is indicative of inflammation; (4) treating (M5) an inflammation (especially chronic) or in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammatory disease, by contacting a tissue having inflammation with an agent that modulates the expression of gene(s) from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for modulating GA; M3 is useful for screening an agent capable of modulating GCA preferably in an inflammation (especially chronic) in a tissue, an allergic detecting an inflammation (especially chronic) in a tissue, an allergic response in a subject, exposure of a subject to a pathogen or sterile inflammation disease (e.g. psoriasis, rheumatoid arthritis, glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal reperfusion injury, ARDS, adult respiratory distress syndrome, inflammatory bowel disease, Crohn's disease, ulcerative colitis, periodontal disease; also bacterial infection, viral infection, M5 is parasitic infection, protozoal infection, fungal infection and M5 is useful for treating one of the above conditions. The present sequence represents a gene differentially expressed in granulocytes. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

Sequence 86574 BP; 22071 A; 20398 C; 21552 G; 22553 T; 0 other;

Query Match	18.6%;	Score 171.6;	DB 24;	Length 86574;
Best Local Similarity	71.9%;	Pred. No. 7.4e-31;		
Matches 299;	Conservative	0;	Mismatches 74;	Indels 43;
				Gaps 4;

QY 428 AACGGAGGAGGAGGAAGGAAGGAGATGATGAGGATGAGGATATATCTC 487
 16224 AGCGAAAGGGAGAGGAGGAGGAGGAAGGATGATGAGGATAAAGGATGCGAAGGGTCTC 16165

QY 488 TGGAGGAGCAAAAGCCCTGTCCAACAAGTCAAAGGCCTGGTGCCCCAGAAAGCAGGCCGACCG 547
||||| | | | ||||| | | | | | | | | | | | | | | |
Db 16164 TGGAGAAGGAGAGACCCTGT --- CAAGTCAAAAAGCGCTGGCGCCCAGAAAGCAGAGGAGCT 16109

	QY	548	TGCGCTAAGAAAAAAAGTGGAAAAAGAACA-----TAGACGAATAATAGACC	593
	Db	16108	TTGCCGAAAAAAAAAGAAAAGAAAAAAACTGGAAAAAGAGAGGAAGTGAGACC	16049

	QY	594	CAGCGCTTAGAGACAAGACCCCTGTGAAAAAAGGCCAAAGCCACAGCCAGCCCAAGAAGCC	653
	Db	16048	TAGTCTTAGAGGCACAGAGCCCTTGGAATATAGGCCAAACACACATTTCAGGCC-----	15998

```

QY      654  AGGATTCAAGAAATGAGAGCCACGCGCTTGGGGGGGACCGGTGCAAAAGTGGGCGCTTCCCTC  713
          |||||
Db 15997 -----CAAGAAATGAGAGCCATGCGCTTGGAGGTCATGGTGCAGAGTGAACCTGCGCCT  159

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[illegible]

QY 774 AGGGGTGTTGCAGGGCAACATGAGAGCCCTCACCCCCAACTTCCTTACG 829
|||||
Db 15893 AGGGGTGTTGTGGGGCAACACTAAAGTCCCTCACCCTCGACTTCAGGTGAGG 15838

RESULT 5	
AAD00297	
ID	AAD00297 standard; DNA; 1481 BP.

AC AAD00297;

DT 05-SEP-2000 (first entry)

Mouse ovary-specific Npm2 gene.

KW oocyte-specific; ovary; Npm2; nucleoplasm; mouse; mammalian ortholog

KW chromosome 14; human chromosome 8p21; contraceptive; gynaecology; cancer;
 KW cell proliferative disorder; cell degenerative disorder; ovulation;
 KW modulator; human infertility; signalling pathway; screen; treatment; ds.
 XX
 OS Mus sp.

Key	Location/Qualifiers
5'UTR	1..150
exon	/*tag= a 151..199
Intron	/*tag= b /number= 1 200..230
misc_feature	/*tag= c /number= 1 215
exon	/*tag= d /note= "Corresponds to 314 missing nucleotides of intron 1 between bases 214 and 216" 231..394
CDS	/*tag= e /number= 2 337..1177
Intron	/*tag= f /product= "Mouse ovary-specific Npm2 protein" /note= "Coding region is interrupted by 8 introns" 395..425
misc_feature	/*tag= g /number= 2 410
exon	/*tag= h /note= "Corresponds to 105 missing nucleotides of intron 2 between bases 409 and 411" 426..511
misc_feature	/*tag= i /number= 3 473
Intron	/*tag= j /note= "The 'T' is replaced with 'G' in the cDNA" 512..542
misc_feature	/*tag= k /number= 3 527
exon	/*tag= l /note= "Corresponds to 63 missing nucleotides of intron 3 between bases 526 and 528" 543..656
Intron	/*tag= m /number= 4 657..687
misc_feature	/*tag= n /number= 4 672
exon	/*tag= o /note= "Corresponds to 2771 (2.77kb) missing nucleotides of intron 4 between bases 671 and 673" 688..781
Intron	/*tag= p /number= 5 782..812
misc_feature	/*tag= q /number= 5 797
exon	/*tag= r /note= "Corresponds to 1321 (1.32kb) missing nucleotides of intron 5 between bases 796 and 798" 813..964
Intron	/*tag= s /number= 6 965..995
misc_feature	/*tag= t /number= 6 980

```

FT      /tag= u
FT      /note= "Corresponds to 157 missing nucleotides of
FT      intron 6 between bases 979 and 981"
FT      996..1036
FT      /tag= v
FT      /number= 7
FT      1037..1067
FT      /tag= w
FT      /number= 7
FT      1052
FT      /tag= x
FT      /note= "Corresponds to 471 missing nucleotides of
FT      intron 7 between bases 1051 and 1053"
FT      1068..1101
FT      /tag= y
FT      /number= 8
FT      1102..1132
FT      /tag= z
FT      /number= 8
FT      1117
FT      /tag= aa
FT      /note= "Corresponds to 63 missing nucleotides of
FT      intron 8 between bases 1116 and 1118"
FT      1133..1381
FT      /tag= ab
FT      /number= 9
FT      1229
FT      /tag= ac
FT      /note= "The 'C' is replaced with 'T' in the cDNA"
FT      1361..1366
FT      /tag= ad
FT      1382..1481
FT      /tag= ae
FT

```

PN	WO2000024755-A1.
XX	
PD	04-MAY-2000.
XX	
PF	28-OCT-1999; 99WO-US25209.
XX	
PR	28-OCT-1998; 98US-0106020.
XX	
PA	(BAYU) BAYLOR COLLEGE MEDICINE
XX	
PI	Matzuk MM, Wang P;
XX	
DR	WPI; 2000-350684/30. P-PsDB; AAY70951.

PT	01-180, 01-184 and 01-236 polypeptides and nucleic acids encoding them.
PT	useful for evaluating potential contraceptives to block ovulation in a
PT	reversible manner -
XX	
PS	Example 5; Fig 13; 54pp; English

CC The present sequence is the mouse ovary-specific Npm2 gene, mapped to
CC the middle of mouse chromosome 14. It shows linkage to D14Mit32, this
CC region being syntenic to human chromosome 8p21. The clone O1-236 cDNA is
CC used to screen and identify Npm2 gene. It is the mammalian ortholog of
CC *Xenopus laevis* nucleoplasmin (Xnpm2) expressed exclusively in the eggs.
CC It provides in vitro and in vivo reagents for studying ovarian
CC development and function. This sequence has gynaecological and
CC contraceptive activity. Agents which modulate O1-180, O1-184 and O1-236
CC may be used to treat cell proliferative or degenerative disorders, this
CC associated with abnormal expression of these ovary specific genes. This
CC ovary-specific sequence can be used as reagents to evaluate potential
CC contraceptives, to block ovulation in a reversible manner. It is also
CC used to screen for genetic mutations in signalling pathways, that is
CC associated with some form of human infertility or gynaecological cancer.
XX
SQ Sequence 1481 BP; 407 A; 379 C; 391 G; 296 T; 8 other;

Query Match

14.28; Score 131.6; DB 21; Length 1481;

PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
PR 08-NOV-2000; 2000US-0246525.
PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246528.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.
PR 08-NOV-2000; 2000US-0246610.
PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
PR 17-NOV-2000; 2000US-0249213.
PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249244.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249264.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251988.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
XX
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Rosen CA, Barash SC, Ruben SM;
XX
DR WPI; 2001-465570/50.
DR P-PSDB; AAM95834.
XX
XX
PT Isolated nucleic acid molecule encoding a reproductive system antigen
PT is used in preventing, treating or ameliorating a medical condition -
XX
XX
PS Claim 1; SEQ ID NO 1805; 1297pp + Sequence Listing; English.
XX
CC The present invention provides the protein and coding sequences of a
CC number of human reproductive system related antigens. These can be used
CC in the prevention and treatment of reproductive system disorders,
CC including cancer. The present sequence is a coding sequence of the
CC invention.
XX
SQ Sequence 416 BP; 96 A; 105 C; 130 G; 80 T; 5 other;

Query Match 7.8%; Score 72.4; DB 22; Length 416;
Best Local Similarity 50.0%; Pred. No. 8.2e-08;
Matches 203; Conservative 1; Mismatches 196; Indels 6; Gaps 1;

QY 51 GGAGGAAAGCAGTCAGCAGCCGTGCTGCGGCTGCGAGCTCAGTCAGAGAGGCGGAC 110

Db 2 GCACGAGAGGGGTGTACTGGCGCGGTAGAGGCTGTGAGCTCTCCGGCCACCCGCTC 61
QY 111 TTGACCTTCAGACCCAGCTGGAGGGGAGAGCAGAGCTGCTGCTTCATACGAT 170
Db 62 CTCACCTTTAAGGTAGAGGAGAGGATGATGCCGASACGTGNCCTGGCACTAACCATGCT 121
QY 171 TTGCTTGGGGGAGAAAGCCAAAGAGAGATGCATCCGCTGAGATCCTGCCCCACCAAA 230
Db 122 CTGCCTCACCAGGAGGCCAAAGACGAGTGAATGTGTTAGAGTTGTGCGCCGGA---- 177
QY 231 CCAGGAGGACAAGAGATGCAGCCGGTCACCAATTGCTCACTCCAGGCCCTCAGTCTCCC 290
Db 178 --ACCATGACCATCAGGAGATCGCAGTCCCTGTGGCCAACTCAAGCTGTCTGCCAACCC 235
QY 291 CATGCTCTCCATGTAAGATGCAGCTTCTCCCGCAGTTACTTCCAGCTCCGGGCTGG 350
Db 236 CATGCTCAGTCTGATGACTTCCAGCTCCAACCACTGTAACTTCCGCCCTGAAGTCGGG 295
QY 351 CTCAGGACCCGCTGTCTCAGTGGCCAGGAGACGTTATGAAGCATCAGACCTAACCTGGGA 410
Db 296 TTCTGGCCCTGTGCGGNTCANTGGCGGCCACCAAGATTGTTACGATGAGCAATGATGTTTC 355
QY 411 GGAGGAGGAGGAAGAGAGGGGAGGAGGAGGAAGGAAGAGAA 456
Db 356 TGAGGAGGAGAGCGAGGAGGAAGGAAGAGACANTGATGAGGAAGA 401

RESULT 7
ABL97097
ID ABL97097 standard; cDNA; 416 BP.

XX ABL97097;

DT 21-JUN-2002 (first entry)

XX Human testicular antigen encoding cDNA SEQ ID NO: 765.

XX Human; testicular antigen; testes; cancer; metastasis; immune disorder;

KW reproductive system disorder; urinary system disorder; gene therapy;

KW cardiovascular disorder; respiratory disorder; neurological disorder;

KW gastrointestinal disease; infection; cytostatic; gene; ss.

XX Homo sapiens.

XX WO200155317-A2.

XX 02-AUG-2001.

XX 17-JAN-2001; 2001WO-US01329.

XX 31-JAN-2000; 2000US-0179065.

XX 04-FEB-2000; 2000US-0180628.

XX 24-FEB-2000; 2000US-0184664.

XX 02-MAR-2000; 2000US-0186350.

XX 16-MAR-2000; 2000US-0189874.

XX 17-MAR-2000; 2000US-0190076.

XX 18-APR-2000; 2000US-0198123.

XX 19-MAY-2000; 2000US-0205515.

XX 07-JUN-2000; 2000US-0209467.

XX 28-JUN-2000; 2000US-0214886.

XX 30-JUN-2000; 2000US-0215135.

XX 07-JUL-2000; 2000US-0216647.

XX 07-JUL-2000; 2000US-0216880.

XX 11-JUL-2000; 2000US-0217487.

XX 11-JUL-2000; 2000US-0217496.

XX 14-JUL-2000; 2000US-0218290.

XX 26-JUL-2000; 2000US-0220963.

XX 26-JUL-2000; 2000US-0220964.

XX 14-AUG-2000; 2000US-0224518.

XX 14-AUG-2000; 2000US-0224519.

XX 14-AUG-2000; 2000US-0225213.

XX 14-AUG-2000; 2000US-0225214.

PR	14-AUG-2000;	2000US-0225266.
PR	14-AUG-2000;	2000US-0225267.
PR	14-AUG-2000;	2000US-0225268.
PR	14-AUG-2000;	2000US-0225270.
PR	14-AUG-2000;	2000US-0225447.
PR	14-AUG-2000;	2000US-0225757.
PR	14-AUG-2000;	2000US-0225758.
PR	14-AUG-2000;	2000US-0225759.
PR	18-AUG-2000;	2000US-0226279.
PR	22-AUG-2000;	2000US-0226681.
PR	22-AUG-2000;	2000US-0226868.
PR	22-AUG-2000;	2000US-0227182.
PR	23-AUG-2000;	2000US-0227009.
PR	30-AUG-2000;	2000US-0228924.
PR	01-SEP-2000;	2000US-0229287.
PR	01-SEP-2000;	2000US-0229343.
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PR	01-SEP-2000;	2000US-0229345.
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PR	05-SEP-2000;	2000US-0229513.
PR	06-SEP-2000;	2000US-0230437.
PR	06-SEP-2000;	2000US-0230438.
PR	08-SEP-2000;	2000US-0231242.
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PR	08-SEP-2000;	2000US-0231414.
PR	08-SEP-2000;	2000US-0232080.
PR	08-SEP-2000;	2000US-0232081.
PR	12-SEP-2000;	2000US-0231968.
PR	14-SEP-2000;	2000US-0232397.
PR	14-SEP-2000;	2000US-0232398.
PR	14-SEP-2000;	2000US-0232399.
PR	14-SEP-2000;	2000US-0232400.
PR	14-SEP-2000;	2000US-0232401.
PR	14-SEP-2000;	2000US-0233063.
PR	14-SEP-2000;	2000US-0233064.
PR	14-SEP-2000;	2000US-0233065.
PR	21-SEP-2000;	2000US-0234223.
PR	21-SEP-2000;	2000US-0234274.
PR	25-SEP-2000;	2000US-0234997.
PR	25-SEP-2000;	2000US-0234998.
PR	26-SEP-2000;	2000US-0235484.
PR	27-SEP-2000;	2000US-0235834.
PR	27-SEP-2000;	2000US-0235836.
PR	29-SEP-2000;	2000US-0236327.
PR	29-SEP-2000;	2000US-0236367.
PR	29-SEP-2000;	2000US-0236368.
PR	29-SEP-2000;	2000US-0236369.
PR	29-SEP-2000;	2000US-0236370.
PR	02-OCT-2000;	2000US-0236802.
PR	02-OCT-2000;	2000US-0237037.
PR	02-OCT-2000;	2000US-0237038.
PR	02-OCT-2000;	2000US-0237039.
PR	02-OCT-2000;	2000US-0237040.
PR	13-OCT-2000;	2000US-0239935.
PR	13-OCT-2000;	2000US-0239937.
PR	20-OCT-2000;	2000US-0241221.
PR	20-OCT-2000;	2000US-0241785.
PR	20-OCT-2000;	2000US-0241786.
PR	20-OCT-2000;	2000US-0241787.
PR	20-OCT-2000;	2000US-0241808.
PR	20-OCT-2000;	2000US-0241809.
PR	20-OCT-2000;	2000US-0241826.
PR	01-NOV-2000;	2000US-0244617.
PR	08-NOV-2000;	2000US-0246474.
PR	08-NOV-2000;	2000US-0246475.
PR	08-NOV-2000;	2000US-0246476.
PR	08-NOV-2000;	2000US-0246477.
PR	08-NOV-2000;	2000US-0246478.
PR	08-NOV-2000;	2000US-0246523.
PR	08-NOV-2000;	2000US-0246524.

PR	08-NOV-2000;	2000US-02465525;
PR	08-NOV-2000;	2000US-02465526;
PR	08-NOV-2000;	2000US-02465527;
PR	08-NOV-2000;	2000US-02465528;
PR	08-NOV-2000;	2000US-02465532;
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PR	08-NOV-2000;	2000US-02466610;
PR	08-NOV-2000;	2000US-02466611;
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PR	17-NOV-2000;	2000US-0249207;
PR	17-NOV-2000;	2000US-0249208;
PR	17-NOV-2000;	2000US-0249209;
PR	17-NOV-2000;	2000US-0249210;
PR	17-NOV-2000;	2000US-0249211;
PR	17-NOV-2000;	2000US-0249212;
PR	17-NOV-2000;	2000US-0249213;
PR	17-NOV-2000;	2000US-0249214;
PR	17-NOV-2000;	2000US-0249215;
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PR	17-NOV-2000;	2000US-0249217;
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PR	17-NOV-2000;	2000US-0249224;
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PR	17-NOV-2000;	2000US-0249300;
PR	01-DEC-2000;	2000US-0250160;
PR	01-DEC-2000;	2000US-0250391;
PR	05-DEC-2000;	2000US-0251030;
PR	05-DEC-2000;	2000US-0251988;
PR	05-DEC-2000;	2000US-0256719;
PR	06-DEC-2000;	2000US-0251476;
PR	08-DEC-2000;	2000US-0251859;
PR	08-DEC-2000;	2000US-0251868;
PR	08-DEC-2000;	2000US-0251869;
PR	08-DEC-2000;	2000US-0251989;
PR	08-DEC-2000;	2000US-0251990;
PR	11-DEC-2000;	2000US-0254097;
PR	05-JAN-2001;	2001US-0259678;

(HUMA-) HUMAN GENOME SCI INC.

Rosen CA, Barash SC, Ruben SM;

WPT; 2001-483232/52.

Nucleic acids encoding 973 human testicular antigen polypeptides, useful for preventing, diagnosing and/or treating testicular cancer

Claim 1; SEQ ID NO 765; 766pp; English.

The present invention provides the protein and coding sequences of 973 human testicular antigens, and fragments of their genomic sequences. The sequences can be used in the treatment of cardiovascular, urinary system, reproductive system, immune, respiratory, neurological and gastrointestinal disorders, infections, and particularly cancer, especially testicular cancers. The present sequence is a cDNA of the invention.

Sequence 416 BP; 96 A; 105 C; 130 G; 80 T; 5 other;

Query Match	7.8%;	Score 72.4;	DB 23;	Length 416;
Best Local Similarity	50.0%;	Pred. No. 8.2e-08;		
Matches 203; Conservative	1;	Mismatches 196;	Indels 6;	Gaps 1;

QY 51 GGAGGAAAAGCGACTGACGACCCTGCTCTGGGGCTGCAGCTCAGTCAGAGAGCCGGAC 110
| | | | | | | | | | | | | | | | | |
db 2 GCACGAGAGGGGTGTACTCTGGGGCGTAAAGAGCTGTGAGCTCTCCGCCACACCCGCTC 61

QY 111 TTGACCTTCAGACCCAGCTTGAGGGGAAACAGAGCTGCAGGCTGTGCTTCATACGAT 170
| | | | | | | | | | | : | | | | |

Db 62 CTTTACCCTTTAAGGTAGAGGAAGAGGATGATGGCGAASACGTGNCCTGGCACTAACCATGCT 121

QY 171 TTGCTTTGGGGAGAAAGCCAAAGAGAGATGCATCGCGTGGAGATCTTGGCCCCAGCAAA 230
111 111 11111111 111 11 11 1 11 111

Db 122 CTGCCTTCACCGAGGAGAGCCAAAGACGAGTGTATGTGGTAGAAGTTGTGGCCCCGGA --- 177

QY 231 CCAGGAGGACAGAAGATGCCAGCCGGTCAACCATTTGCCACTCCAGGCGCTAGTCTCC 290
1 111 11 1 111 1 111 1 11 11 11 11 11

Db 178 -- ACCATGACCATCAGAGAGATCGCAGTCCCTGTGGCCAACTCAAGCTGTCTGCCAAC 235

QY 291 CATGGTCTCCATGCTTAGGAGTGCAGCTTTTCCGCCAGTTACTTTCCAGCTCCGGGCTGG 350
1111 11 111 1 1111 11 11 11 1111 11 1 11 11

Db 236 CATGCTCAGTCTGGATGACTTCCAGCTCCAACCACTGTAAACCTTCCGCCCTGAAGTCGG 295

QY 351 CTCAGGACCCGTGTTCTCTCAGTGGGCCAGSAAAGTTATGAAGCATTCAGACCTAACCTGGGA 410
11 11 11 111 111 111 1 1 11 1

Db 296 TTCTGGCCCTGTCCGNTCANTGGGCGCGCACAGATTGTTACGATGAGCAATGATGTTTC 355

QY 411 GGAGGAGGAGGAAGAAGAGGGGAGAGAGAGGAAGAAGAGAGAA 456
11111111 11 111 111 1111 11 11 11 111

Db 356 TGAGGAGGAGAGCGAGGAAGAGAGGAAGAGACANTGATGAGCAAGAA 401

```

RESULT 8
AAZ23891
ID AAZ23891 standard; DNA; 49999 BP.
XX
XX
AC AAZ23891;
XX
XX 25-JAN-2000 (first entry)
DT
DE Murine LOBO genomic DNA fragment 1.
XX
XX LOBO; long bones; bone development; bone extension; skull; osteopathic;
KW diagnostic; pharmaceutical; gene therapy; transgenic animal; disease;
XX spondyloepiphyseal dysplasia; achondroplasia; murine; ds.
OS
XX Mus musculus.
XX
XX WO9950284-A2.
XX
XX 07-OCT-1999.
XX
XX 26-MAR-1999; 99WO-EP02055.
XX
XX 27-MAR-1998; 98DE-1013799.
XX
XX (ROSE/) ROSENTHAL A.
XX
XX Rosenthal A, Rump A, Hess J, Aigner T, Wirth T;
XX
XX WPI; 1999-601320/51.
XX
XX Nucleic acids encoding proteins which influence bone development,
XX useful for treating and studying bone disorders -
XX
XX Example 3; Page 69-97; 391pp; German.
XX
XX This invention describes novel nucleic acids (I; designated LOBO (long
XX bones)) encoding proteins influencing bone development in mammals. The
XX proteins of the invention reduce and/or inactivate bone extension (i.e.
XX development), with exception of the skull and have osteopathic activity
XX The nucleic acid molecules, proteins and antibodies can be used in
XX diagnostic or pharmaceutical compounds e.g. for gene therapy. The method
XX and nucleic acid molecules, etc. are useful for production of transgenic
XX animals, especially a transgenic mouse for the study of diseases
XX associated with bone development, e.g. spondyloepiphyseal dysplasia and
XX achondroplasia. This sequence encodes the murine LOBO protein described
XX in the method of the invention.
XX
XX Sequence 49999 BP; 13210 A; 11814 C; 10825 G; 14150 T; 0 other;
XX

```

[illegible]

RESULT 9
 AA223896
 ID AA223896 standard; DNA; 49999 BP.
 AC AA223896;
 DT 25-JAN-2000 (first entry)
 DE Murine LOBO homologue genomic DNA fragment 2.
 KW LOBO; long bones; bone development; bone extension; skull; osteopathic;
 KW diagnostic; pharmaceutical; gene therapy; transgenic animal; disease;
 KW spondyloepiphyseal dysplasia; achondroplasia; murine; ds.
 OS Mus musculus.
 PN WO9950284-A2.
 PD 07-OCT-1999.
 PF 26-MAR-1999; 99WO-EP02055.
 PR 27-MAR-1998; 98DE-1013799.
 PA (ROSE/) ROSENTHAL A.
 PI Rosenthal A, Rump A, Hess J, Aigner T, Wirth T;
 DR WPI; 1999-601320/51.
 PT Nucleic acids encoding proteins which influence bone development,
 PR useful for treating and studying bone disorders -
 PS
 PS Example 3; Page 161-189; 391pp; German.
 CC This invention describes novel nucleic acids (I; designated LOBO (long
 CC bones)) encoding proteins influencing bone development in mammals. The
 CC proteins of the invention reduce and/or inactivate bone extension (i.e.
 CC development), with exception of the skull and have osteopathic activity.
 CC The nucleic acid molecules, proteins and antibodies can be used in
 CC diagnostic or pharmaceutical compounds e.g. for gene therapy. The methods
 CC and nucleic acid molecules, etc. are useful for production of transgenic
 CC animals, especially a transgenic mouse for the study of diseases
 CC associated with bone development, e.g. spondyloepiphyseal dysplasia and
 CC achondroplasia. This sequence encodes the murine LOBO protein described
 CC in the method of the invention.
 CC
 CC Sequence 49999 BP; 13135 A; 11787 C; 10868 G; 14209 T; 0 other:

PR	08-NOV-2000;	2000US-0246609.
PR	08-NOV-2000;	2000US-0246610.
PR	08-NOV-2000;	2000US-0246611.
PR	08-NOV-2000;	2000US-0246613.
PR	17-NOV-2000;	2000US-0249207.
PR	17-NOV-2000;	2000US-0249208.
PR	17-NOV-2000;	2000US-0249209.
PR	17-NOV-2000;	2000US-0249210.
PR	17-NOV-2000;	2000US-0249211.
PR	17-NOV-2000;	2000US-0249212.
PR	17-NOV-2000;	2000US-0249213.
PR	17-NOV-2000;	2000US-0249214.
PR	17-NOV-2000;	2000US-0249215.
PR	17-NOV-2000;	2000US-0249216.
PR	17-NOV-2000;	2000US-0249217.
PR	17-NOV-2000;	2000US-0249218.
PR	17-NOV-2000;	2000US-0249244.
PR	17-NOV-2000;	2000US-0249245.
PR	17-NOV-2000;	2000US-0249264.
PR	17-NOV-2000;	2000US-0249265.
PR	17-NOV-2000;	2000US-0249297.
PR	17-NOV-2000;	2000US-0249299.
PR	17-NOV-2000;	2000US-0249300.
PR	01-DEC-2000;	2000US-0250160.
PR	01-DEC-2000;	2000US-0250391.
PR	05-DEC-2000;	2000US-0251030.
PR	05-DEC-2000;	2000US-0251988.
PR	05-DEC-2000;	2000US-0256719.
PR	06-DEC-2000;	2000US-0251479.
PR	08-DEC-2000;	2000US-0251856.
PR	08-DEC-2000;	2000US-0251868.
PR	08-DEC-2000;	2000US-0251869.
PR	08-DEC-2000;	2000US-0251989.
PR	08-DEC-2000;	2000US-0251990.
PR	11-DEC-2000;	2000US-0254097.
PR	05-JAN-2001;	2001US-0259678.

PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX
PI Rosen CA, Barash SC, Ruben SM;
XX
XX
DR WPT; 2001-483426/52.
XX
PT Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and
PT metastasis -

PS Disclosure; SEQ ID NO 21883; 3071pp + Sequence Listing; English.

CC AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytosolic
CC activity, and can be used in gene therapy and vaccine production. (I)
CC proteins and polynucleotides may be used in the prevention, diagnosis and
CC treatment of diseases associated with inappropriate (I) expression. For
CC example, they may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patient's genome
CC that affect the activity of (I) by expressing inactive proteins or to
CC supplement the patients own production of (I). Additionally, (I)
CC polynucleotides may be used to produce the secreted (I), by inserting
CC the nucleic acids into a host cell and culturing the cell to express the
CC protein. (I) proteins and polynucleotides may be used to prevent,
CC diagnose and treat immune/haematopoietic-related diseases, especially
CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
CC to AAK87694 represent human immune/haematopoietic antigen genomic
CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
CC represent sequences used in the exemplification of the present invention.

50 Sequence 33923 BP; 9925 A; 8140 C; 7946 G; 7912 T; 0 other;

Query Match	7.58;	Score 69.2;	DB 22;	length 33923;
Best Local Similarity	58.18;	Pred. No. 2e-06;		
Matches 122;	Conservative	0;	Mismatches 88;	Indels 0;

[illegible]

```

RESULT 11
AAS90676
ID AAS90676 standard; cDNA; 341 bp
....

```

AC AAS90676;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #26480.

KN Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss

OS Homo sapiens.

PN WO200175067-A2

PD 11-OCT-2001

30-MAR-2001; 2001WO-US08631

PR	31-MAR-2000; 2000US-0540217
PR	23-AUG-2000; 2000US-0649167

PA (HYSE-) HYSEQ INC

PI Drmanac RT, Liu C, Tang YT,

DR WPI; 2001-639362/73.

XX
SE

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -

PS Claim 1; SEQ ID NO 26480; 103pp; English

The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal activity of (II) or to treat disease states involving (II). (II) is useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic coding sequences of the invention.

AC AAS68926;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #4730.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI; 2001-639362/73.
DR P-PSDB; ABG04739.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
PS Claim 1; SEQ ID NO 4730; 103pp; English.
XX
CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 291 BP; 129 A; 22 C; 112 G; 28 T; 0 other;
XX
Query Match 7.2%; Score 66.4; DB 23; Length 291;
Best Local Similarity 54.0%; Pred. No. 2e-06;
Matches 136; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

QY 362 TGTTCCTCAGTGGCCAGCAAGCTTATGAAGCATCAGACCTAACCCTGGAGGAGGAGG 421
DB 8 TTTGCCATATTGGCCAGGCTAGTCTCGAAGCTCTGACCTCAGGGGAGGAGGAGGAGG 67
QY 422 AAGAGAAGGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 481
DB 68 AGGAGGAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 127
QY 482 TATCTCTGAGGAGCAAAAGCCCTCAACAAGTCAAAAGGCTGTGCCCCAGAGCAGG 541
DB 128 AAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 187

QY 542 CGAGCGTGCTAAGAAAAAAGCTGGAAAAAGAGAGAGGAATTAAGCCCGCTTA 601
DB 188 AAGAAG 247
QY 602 GAGACAGAGAGCC 613
DB 248 GAGATCAATATCC 259
RESULT 14
AAS68538
ID AAS68538 standard; cDNA; 1680 BP.
XX
AC AAS68538;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #4342.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI; 2001-639362/73.
DR P-PSDB; ABG04351.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
PS Claim 1; SEQ ID NO 4342; 103pp; English.
XX
CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 1680 BP; 574 A; 286 C; 485 G; 335 T; 0 other;
XX
Query Match 7.1%; Score 65.8; DB 23; Length 1680;

•
•
•
•

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: January 24, 2003, 03:53:00 ; Search time 2237 Seconds
(without alignments)
6689.598 Million cell updates/sec

Title: US-09-844-864-16

Perfect score: 924

Sequence: 1 cagccgcgtctctgcgcg.....tttgcgcgcagcttatg 924

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

EST: *
1: em_estba: *
2: em_esthum: *
3: em_estlin: *
4: em_estmu: *
5: em_estov: *
6: em_estpl: *
7: em_estro: *
8: em_hlc: *
9: gb_est1: *
10: gb_est2: *
11: gb_hlc: *
12: gb_est3: *
13: gb_est4: *
14: gb_est5: *
15: em_estfun: *
16: em_estom: *
17: gb_gss: *
18: em_gss_hum: *
19: em_gss_inv: *
20: em_gss_pln: *
21: em_gss_vrt: *
22: em_gss_fun: *
23: em_gss_mam: *
24: em_gss_mus: *
25: em_gss_other: *
26: em_gss_pro: *
27: em_gss_rod: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	517	56.0	517	9	AI391464	AI391464 tf96g05.x
2	455	49.2	455	9	AI016313	AI016313 ot78a10.s
3	433	46.9	433	13	BM055230	BM055230 ie92b10.y
4	432.2	46.8	451	13	BM054976	BM054976 ie92b10.x
5	418.6	45.3	1324	14	BM809804	BM809804 AGENCOURT
6	414.4	44.8	854	14	BQ878692	BQ878692 AGENCOURT

7	410.6	44.4	915	14	BQ684634	BQ684634 AGENCOURT
8	405.8	43.9	872	14	BQ878312	BQ878312 AGENCOURT
9	405	43.8	405	12	BF594409	BF594409 7105a05.x
10	397.4	43.0	399	10	AW731946	AW731946 ba03e06.x
11	389.6	42.2	935	14	BQ684424	BQ684424 AGENCOURT
12	378.4	41.0	966	14	BQ682257	BQ682257 AGENCOURT
13	357.8	38.7	361	10	AW002370	AW002370 wa61a07.x
14	357.8	38.7	361	12	BF592761	BF592761 7194b04.x
15	357.8	38.7	361	12	BF057162	BF057162 7k16g07.x
16	352.4	38.1	377	12	BF804737	BF804737 CMO-CT009
17	350.6	38.0	378	12	BF223902	BF223902 7q36b08.x
18	350.6	37.9	1047	14	BQ888606	BQ888606 AGENCOURT
19	280.4	30.3	491	13	BI913169	BI913169 603179964
20	234.2	25.3	636	10	BB559308	BB559308 BB559308
21	178.4	19.3	580	13	BI439005	BI439005 1c25d05.y
22	158.6	17.2	280	10	BB284820	BB284820 BB284820
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24	103.2	11.2	548	12	BG515101	BG515101 da659h10.
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26	103.2	11.2	658	9	AL782493	AL782493 AL782493
27	102	11.0	566	9	AL645040	AL645040 AL645040
28	96.8	10.5	593	10	BE026525	BE026525 db227f07.x
29	96.4	10.4	574	13	BJ094747	BJ094747 BJ094747
30	96.4	10.4	600	14	BQ387680	BQ387680 NISC_mn25
31	96.4	10.4	645	13	BJ098416	BJ098416 BJ098416
32	96	10.4	590	13	BI941416	BI941416 df88a05.y
33	95.8	10.4	555	12	BG021389	BG021389 df87a05.x
34	94.4	10.2	380	10	BE506675	BE506675 db87d01.y
35	94.2	10.2	522	12	BG886286	BG886286 da651d02.
36	94.2	10.2	551	12	BG016384	BG016384 df88a05.x
37	93.8	10.2	594	12	BF427228	BF427228 df87a05.y
38	93.4	10.1	564	10	BE678406	BE678406 df55d03.x
39	91.4	9.9	531	10	BE192089	BE192089 db87d01.x
40	90.2	9.8	460	12	BG515848	BG515848 da659h10.
41	89	9.6	593	10	BE026442	BE026442 db26f09.x
42	87.2	9.4	554	10	BE680412	BE680412 df79b03.y
43	84.8	9.2	541	9	AL595075	AL595075 AL595075
44	83.6	9.0	549	14	BQ387679	BQ387679 NISC_mn25
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ALIGNMENTS

RESULT 1
AI391464/c 517 bp mRNA linear EST 02-FEB-1999
LOCUS tf96g05.x1 NCI-CGAP-CLL1 Homo sapiens CDNA clone IMAGE:2107160 3'
DEFINITION similar to contains element MER22 repetitive element ;, mRNA
sequence.

ACCESSION AI391464
VERSION AI391464
KEYWORDS AI391464.1 GI:4217468
SOURCE EST.
ORGANISM human.

REFERENCE
AUTHORS Homo sapiens
TITLE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

JOURNAL
COMMENT NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
Tissue Procurement: Ash Alizadeh, John Byrd, M.D., Mike Grever,
M.D., Louis M. Staudt, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/dbp/image/image.html
Seq primer: -400p from Gibco

BM055230
LOCUS 433 bp mRNA linear EST 12-MAR-2002
DEFINITION Ie92b10.y1 Melton Normalized Human Islet 4 N4-HIS 1 Homo sapiens
ACCESSION BM055230
VERSION BM055230.1 GI:16812965
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 433)
AUTHORS Melton,D., Brown,J., Kenty,G., Permutt,A., Lee,C., Kaestner,K.,
Lemishka,I., Scearce,M., Brestelli,J., Gradwohl,G., Clifton,S.,
Hillier,L., Marra,M., Pape,D., Wylie,T., Martin,J., Blistain,A.,
Schmitt,A., Theising,B., Ritter,E., Ronko,I., Bennett,J., Cardenas
, M., Gibbons,M., McCann,R., Cole,R., Tsagarishvili,R., Williams,T.,
Jackson,Y. and Bowers,Y.
Endocrine Pancreas Consortium
Unpublished (2000)
Other_ESTs: Ie92b10.x1
Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
Endocrine Pancreas Consortium
Harvard University, Howard Hughes Medical Institute
Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
MA 02138
Tel: 617-495-1812
Fax: 617-495-8557
Email: dmelton@biohp.harvard.edu
Library was constructed by Dr. Douglas Melton DNA sequencing by:
Washington University Genome Sequencing Center For information on
obtaining a clone please contact: Juliana Brown
(brown@fas.harvard.edu) This sequence now available from the IMAGE
consortium, for clone orders contact: info@image.llnl.gov.

FEATURES

Location/Qualifiers

1..433

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:5674315"

/clone_lib="Melton Normalized Human Islet 4 N4-HIS 1"

/sex="Both"

/tissue_type="Islets of Langerhans"

/dev_stage="Adult"

/lab_host="DH10B"

/note="Organ: Pancreas; Vector: pSPORT1; Site_1: Not 1;
Site_2: Sal 1; Starting library constructed using
Superscript Plasmid Library kit (Life Technologies). cDNA
made by oligo-dT priming. Size-selected by column
fractionation; average insert size 1.08 kb. Library was
amplified once on solid support and plasmid DNA from
library was prepared. The library DNA was normalized by
method #4 from Bonaldo, Lennon, and Soares 1996 Genome
Research 6:791-806; 0.5 microgram single-stranded library
plasmid DNA was mixed with 5 micrograms PCR product
representing library inserts and hybridized to an Ecot of
20. Single-stranded (unhybridized) plasmids were isolated
by hydroxyapatite chromatography and used to make this
library."

BASE COUNT 137 a 98 c 143 g 55 t
ORIGIN

Query Match

Best Local Similarity 100.0%; Score 433; DB 13; Length 433;
Matches 433; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 391 GCATCAGACCTTAACCTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAA 450
DB 1 GCATCAGACCTTAACCTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAA 60
OY 451 GAGCAGATGATGAGATGAGATGAGATGATATCTCTGAGAGAGCAAGCCCTGTCAA 510
DB 61 GAGCAGATGATGAGATGAGATGAGATGATATCTCTGAGAGAGCAAGCCCTGTCAA 120

OY 511 CAAGTCAAAAGCGTGGTCCCGCAGAAAGCAGCGCGTGGCTTAAGAAAAAAGCTGAA 570
DB 121 CAAGTCAAAAGCGTGGTCCCGCAGAAAGCAGCGCGTGGCTTAAGAAAAAAGCTGAA 180
OY 571 AAAGAAGAGAGAGAAATTAAGAGCCAGCGTTAGAGACAAAGAGCCCTGTGAAGAGCCCAA 630
DB 181 AAAGAAGAGAGAGAAATTAAGAGCCAGCGTTAGAGACAAAGAGCCCTGTGAAGAGCCCAA 240
OY 631 GCCACAGCCAGAGCCCAAGAGCCAGATTCAGAAATGAGAGAGCCAGCCCTTGGGGGCA 690
DB 241 GCCACAGCCAGAGCCCAAGAGCCAGATTCAGAAATGAGAGAGCCAGCCCTTGGGGGCA 300
OY 691 CGGTGCAAAAGTGGGCGCTTCCCTGGGCTGTGCTGCAGGACAGAGGTCCTGTCAGCCC 750
DB 301 CGGTGCAAAAGTGGGCGCTTCCCTGGGCTGTGCTGCAGGACAGAGGTCCTGTCAGCCC 360
OY 751 CTCACACCTGTGTCTGAATGCAACAGGGGTGTGGGGGGCAACATGAGAGCCCTCACCC 810
DB 361 CTCACACCTGTGTCTGAATGCAACAGGGGTGTGGGGGGCAACATGAGAGCCCTCACCC 420
OY 811 CCAACTCTCCACT 823
DB 421 CCAACTCTCCACT 433

RESULT 4

BM054976/c

LOCUS 451 bp mRNA linear EST 12-MAR-2002
DEFINITION Ie92b10.x1 Melton Normalized Human Islet 4 N4-HIS 1 Homo sapiens
ACCESSION BM054976
VERSION BM054976
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 451)
AUTHORS Melton,D., Brown,J., Kenty,G., Permutt,A., Lee,C., Kaestner,K.,
Lemishka,I., Scearce,M., Brestelli,J., Gradwohl,G., Clifton,S.,
Hillier,L., Marra,M., Pape,D., Wylie,T., Martin,J., Blistain,A.,
Schmitt,A., Theising,B., Ritter,E., Ronko,I., Bennett,J., Cardenas
, M., Gibbons,M., McCann,R., Cole,R., Tsagarishvili,R., Williams,T.,
Jackson,Y. and Bowers,Y.
Endocrine Pancreas Consortium
Unpublished (2000)
Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
Endocrine Pancreas Consortium
Harvard University, Howard Hughes Medical Institute
Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
MA 02138
Tel: 617-495-1812
Fax: 617-495-8557
Email: dmelton@biohp.harvard.edu
Library was constructed by Dr. Douglas Melton DNA sequencing by:
Washington University Genome Sequencing Center For information on
obtaining a clone please contact: Juliana Brown
(brown@fas.harvard.edu) This sequence now available from the IMAGE
consortium, for clone orders contact: info@image.llnl.gov.

FEATURES

Location/Qualifiers
1..451
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5674315"
/clone_lib="Melton Normalized Human Islet 4 N4-HIS 1"
/sex="Both"
/tissue_type="Islets of Langerhans"
/dev_stage="Adult"
/lab_host="DH10B"
/note="Organ: Pancreas; Vector: pSPORT1; Site_1: Not 1;
Site_2: Sal 1; Starting library constructed using
Superscript Plasmid Library kit (Life Technologies). cDNA
made by oligo-dT priming. Size-selected by column

fractionation; average insert size 1.08 kb. Library was amplified once on solid support and plasmid DNA from library was prepared. The library DNA was normalized by method #4 from Bonaldo, Lennon, and Soares 1996 Genome Research 6:791-806; 0.5 microgram single-stranded library plasmid DNA was mixed with 5 micrograms PCR product representing library inserts and hybridized to an Ecot of 20. Single-stranded (unhybridized) plasmids were isolated by hydroxyapatite chromatography and used to make this library."

BASE COUNT 61 a 136 c 110 g 144 t
ORIGIN

Query Match 46.8%; Score 432.2; DB 13; Length 451;
Best Local Similarity 98.9%; Pred. No. 8.9e-63;
Matches 446; Conservative 0; Mismatches 3; Indels 2; Gaps 1;

QY 445 GAGGAAGAGAGATGATGAGATGAGATGATATATCTCTGAGAGCAAGCCCT 504
DB 451 GAGGAAGAGAGATGATGAGATGAGATGATATATCTCTGAGAGCAAGCCCT 392
QY 505 GTCAACAAGTCAAAAGGCTGCTCCCGAGAGCAGCGCTGCTAAAGAAAAAG 564
DB 391 GTCAACAAGTCAAAAGGCTGCTCCCGAGAGCAGCGCTGCTAAAGAAAAAG 332
QY 565 CTGGAAG 624
DB 331 CTGGAAG 272
QY 625 GCCAAG 682
DB 271 GCCAAG 212
QY 683 GGGGGGACGCTGCAAAAGTGGGCTTCCCTGGGCTGCTGCAAGGACAGGGTGCCTG 742
DB 211 GAGGGGACGCTGCAAAAGTGGGCTTCCCTGGGCTGCTGCAAGGACAGGGTGCCTG 152
QY 743 TCCAGCCCTCCACTCTCTGATGCAACAGAGGCTGCTGCGGGGCAACATGAGAGCC 802
DB 151 TCCAGCCCTCCACTCTCTGATGCAACAGAGGCTGCTGCGGGGCAACATGAGAGCC 92
QY 803 CCTCACCCTCCACTCTCTGATGCAAGAGGCCCCAGTGAAGAGCCCCAGCTGGGGTCA 862
DB 91 CCTCACCCTCCACTCTCTGATGCAAGAGGCCCCAGTGAAGAGCCCCAGCTGGGGTCA 32
QY 863 CAATAAAGTGTGCTGTCAGGAAAAAAA 893
DB 31 CAATAAAGTGTGCTGTCAGGAAAAAAA 1

RESULT 5
BM809804 1324 bp mRNA linear EST 05-MAR-2002
LOCUS AGENCOURT_6581147 NIH_MGC_98 Homo sapiens cDNA clone IMAGE:5454702
DEFINITION 5', mRNA sequence.
ACCESSION BM809804
VERSION BM809804.1 GI:19126627
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 1324)
AUTHORS NIH-MGC http://mgi.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs-rt@mail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LICM1950 row: e column: 07
High quality sequence stop: 357.
location/Qualifiers

FEATURES
source

1. 1324
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5454702"
/clone_lib="NIH_MGC_98"
/tissue_type="astrocytoma grade IV, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: brain; Vector: pOTB7; Site_1: XhoI; Site_2:
ECORI; cDNA made by oligo-dT priming. Directionally
cloned into EcorI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Library constructed by Ling Hong
in the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."

BASE COUNT 383 a 90 c 706 g 60 t 85 others
ORIGIN

Query Match 45.3%; Score 418.6; DB 14; Length 1324;
Best Local Similarity 78.4%; Pred. No. 1.1e-60;
Matches 494; Conservative 0; Mismatches 134; Indels 2; Gaps 2;

QY 106 CGGACTTGACCTTCACACCCAGCTGGAGGGGAAGCAGAGCTGCAGGCTGTGCTTCAT 165
DB 1 CGGACTTGACCTTCACACCCAGCTGGAGGGGAAGCAGAGCTGCAGGCTGTGCTTCAT 60
QY 166 ACCATTGCTTGGGGGAGAAAGCCAAAGAGAGATGCATCGCGTGGAGATCCTGCCCCA 225
DB 61 ACCATTGCTTGGGGGAGAAAGCCAAAGAGAGATGCATCGCGTGGAGATCCTGCCCCA 120
QY 226 GCAAAACAGAGAGAGAGAGATGACGCCGCTCAACATTGCTCACTCCAGGCTCAGTC 285
DB 121 GCAAAACAGAGAGAGAGAGATGACGCCGCTCAACATTGCTCACTCCAGGCTCAGTC 180
QY 286 CTCCCATGCTCTCCATGTAAGAGTGCAGCTTCTCCCCAGTTACTTTCCAGCTCCGG 345
DB 181 CTCCCATGCTCTCCATGTAAGAGTGCAGCTTCTCCCCAGTTACTTTCCAGCTCCGG 240
QY 346 GCTGCTCAGAGACCGCTGCTCAGTGGCCAGAGCGTTATGAGCATCAGACCTAAC 405
DB 241 GCTGCTCAGAGACCGCTGCTCAGTGGCCAGAGCGTTATGAGCATCAGACCTAAC 300
QY 406 TGGATGATGAG 465
DB 301 TGGATGATGAG 360
QY 466 GATGAGGATGATATATCTCTGAGAGAGCAAGCCCTGTCAAAACAAAGTCAAAAGGCTG 525
DB 361 GATGAGGATGATATATCTCTGAGAGAGCAAGCCCTGTCAAAACAAAGTCAAAAGGCTG 419
QY 526 GTGCCCCAG 585
DB 420 GTGCCCCAG 479
QY 586 ATAAG-AGCCAGCTTAGAGACCAAGCCCTGTGAAGAAAGGCCCAAGCCACAGCCAGAGC 644
DB 480 ATAAG-AGCCAGCTTAGAGACCAAGCCCTGTGAAGAAAGGCCCAAGCCACAGCCAGAGC 539
QY 645 CAAGAAGCAGAGATCAAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 704
DB 540 CAAGAAGCAGAGATCAAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 599
QY 705 CTTTCCCTGGGCTGTGCTGACAGGCAAGGG 734
DB 600 CTTTCCCTGGGCTGTGCTGACAGGCAAGGG 629

RESULT 6

BQ878692
 LOCUS BQ878692 854 bp mRNA linear EST 16-AUG-2002
 DEFINITION AGENCOURT_8072876 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6087706
 5', mRNA sequence.
 ACCESSION BQ878692
 VERSION BQ878692.1 GI:22270700
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 854)
 AUTHORS NIH-MGC <http://mhc.ncl.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-remail.nih.gov
 Tissue Procurement: DCTD/DTP
 CDNA Library Preparation: Rubln Laboratory
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: LLCM2324 row: 1 column: 11
 High quality sequence stop: 560.
 Location/Qualifiers
 1..854
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:6087706"
 /clone_11b="NIH_MGC_112"
 /tissue_type="melanotic melanoma, cell line"
 /lab_host="DH10B (phage-resistant)"
 /note="Organ: skin; Vector: pOTB7; Site_1: XhoI; Site_2:
 EcoRI; cDNA made by oligo-dT priming. Directionally cloned
 into EcoRI/XhoI sites using the following 5' adaptor:
 GGCACGAG(G). Library constructed by Ling Hong in the
 laboratory of Gerald M. Rubin (University of California,
 Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
 Superscript II RT (Life Technologies). Note: this is a
 NIH_MGC library."

BASE COUNT 181 a 247 c 253 g 151 t 22 others
 ORIGIN

Query Match 44.8%; Score 414.4; DB 14; Length 854;
 Best Local Similarity 76.5%; Pred. No. 6.7e-60;
 Matches 595; Conservative 0; Mismatches 16; Indels 167; Gaps 1;

OY 1 CAGCCCGCTTCTGCCCCGAGCCATGAATCTCAGTAGCGCCAGTACGAGGAAAG 60
 |||||||
 Db 207 CAGCCCGCTTCTGCCCCGAGCCATGAATCTCAGTAGCGCCAGTACGAGGAAAG 266
 |||||||
 OY 61 GCAGTGACGACCGTCTGCGGCTGCGAGCTCAGTCAGAGAGGCGGACTTGGACCTTC 120
 |||||||
 Db 267 GCAGTGACNMCCGCTCTGCGGCTGCGAGCTCAGTCAGAGAGGCGGACTTGGACCTTC 326
 |||||||
 OY 121 AGACCCAGCTGAGGCGGAGAGCAGAGCTGAGGCTGTGCTTCATACGATTGCTTGGGG 180
 |||||||
 Db 327 AGACCCAGCTGAGGCGGAGAGCAGAGCTGAGGCTGTGCTTCATACGATTGCTTGGGG 386
 |||||||
 OY 181 GAGAAAGCCAAAGAGAGATGATCGCGCTGAGATCCTGCCCCAGCAAAACAGAGAGAC 240
 |||||||
 Db 387 GAGAAAGCCAAAGAGAGATGATCGCGCTGAGATCCTGCCCCAGCAAAACAGAGAGAC 446
 |||||||
 OY 241 AAGAAGATGACGCGGTCACCATTCCTCCTCAGGCGCTCCTCCCATGCTCTCC 300
 |||||||
 Db 447 AAGAAGATGACGCGGTCACCATTCCTCCTCAGGCGCTCCTCCCATGCTCTCC 506
 |||||||
 OY 301 ATGGTAGAGTGAAGCTTCTCCCGAGTTACTTTCAGCTCCGGGCTGGCTCAGAGACC 360
 |||||||
 Db 507 ATGGTAGAGTGAAGCTTCTCCCGAGTTACTTTCAGCTCCNNNGCTGGCTCAGAGACC 566
 |||||||

OY 361 GTGTTCTCAGTGGCCAGGACGTTATGAAGCATCAGACCTAACCTGGAGAGAGAG 420
 |||||||
 Db 567 GTGTTCTCAGTGGCCAGGACGTTAT----- 593
 |||||||
 OY 421 GAAGAAGAGGGGAGAGAGAGAGAGAGAGATGATGAGATGAGATGACAGAT 480
 |||||||
 Db 594 ----- 593
 |||||||
 OY 481 ATATCTTGAGAGAGCAAGCCCTGTCAACACAGTCAAAAGGCTGTGCCCCAGAGCAG 540
 |||||||
 Db 594 ----- 593
 |||||||
 OY 541 GCGAGCGTGGCTAAGAAAAAAGCTGGAAGAAAGAGAGAAATAGAGCCAGCGTT 600
 |||||||
 Db 594 -----GAAGAAAAAGCTGGAAGAAAGAGAGAAATAGAGCCAGCGTT 639
 |||||||
 OY 601 AGAGACAAGAGCCCTGTGAAGAAAGGCCAAGCCAGCCAGAGCCAGAGATTC 660
 |||||||
 Db 640 AGAGACAAGAGCCCTGTGAAGAAAGGCTTNNNCTCAGCCAGAGCCAGAGATTC 699
 |||||||
 OY 661 AAGAATGAGAGAGCCAGCGCTTGGGGGCGACGCTGCAAGTGGGCTTCCCTGGCTGTG 720
 |||||||
 Db 700 AAGAATGAGAGAGCCAGCGCTTGGGGGCGACGCTGCAAGTGGGCTTCCCTGGCTGTG 759
 |||||||
 OY 721 CTGCAAGCAGAGGCTGCTGTCAGAGCCCTCAGCTGTGTGAATGCAACAGGG 778
 |||||||
 Db 760 CTGCAAGCAGAGGCTGCTGTCAGAGCCCTCAGCTGTGTGAATGCAACAGGG 817
 |||||||

RESULT 7
 BQ684634 915 bp mRNA linear EST 15-JUL-2002
 LOCUS BQ684634
 DEFINITION AGENCOURT_8032849 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6091012
 5', mRNA sequence.
 ACCESSION BQ684634
 VERSION BQ684634.1 GI:21797313
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 915)
 AUTHORS NIH-MGC <http://mhc.ncl.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-remail.nih.gov
 Tissue Procurement: DCTD/DTP
 CDNA Library Preparation: Rubln Laboratory
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: LLCM2333 row: f column: 05
 High quality sequence stop: 652.
 Location/Qualifiers
 1..915
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:6091012"
 /clone_11b="NIH_MGC_112"
 /tissue_type="melanotic melanoma, cell line"
 /lab_host="DH10B (phage-resistant)"
 /note="Organ: skin; Vector: pOTB7; Site_1: XhoI; Site_2:
 EcoRI; cDNA made by oligo-dT priming. Directionally cloned
 into EcoRI/XhoI sites using the following 5' adaptor:
 GGCACGAG(G). Library constructed by Ling Hong in the
 laboratory of Gerald M. Rubin (University of California,
 Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
 Superscript II RT (Life Technologies). Note: this is a
 NIH_MGC library."

BASE COUNT 197 a 283 c 277 g 153 t 5 others

ORIGIN

Query Match 44.4%; Score 410.6; DB 14; Length 915;
Best Local Similarity 77.0%; Pred. No. 2.8e-59;
Matches 631; Conservative 0; Mismatches 17; Indels 172; Gaps 4;

QY 1 CAGCCCGCTTCTCTGCGGAGCCATGATCTCAGTAGCCGAGTAGCAGGAGGAAAG 60
DB 207 CAGCCCGCTTCTCTGCGGAGCCATGATCTCAGTAGCCGAGTAGCAGGAGGAAAG 266
QY 61 GCAGTGACGACCGTGTCTGGGGCTGCGAGCTCAGTCAGGAGAGGCGGACTTGACCTTC 120
DB 267 GCAGTGACGACCGTGTCTGGGGCTGCGAGCTCAGTCAGGAGAGGCGGACTTGACCTTC 326
QY 121 AGACCCCACTGGAGGGGAGAGAGAGCTGCAGGCTGTTCCTTATACGATTTGCTTGGGG 180
DB 327 AGACCCCACTGGAGGGGAGAGAGAGCTGCAGGCTGTTCCTTATACGATTTGCTTGGGG 386
QY 181 GAGAAAGCCAAAGAGAGATGATTCGCGTGGAGATCCTGCCCCAGCAAAACCAGAGGAC 240
DB 387 GAGAAAGCCAAAGAGAGATGATTCGCGTGGAGATCCTGCCCCAGCAAAACCAGAGGAC 446
QY 241 AAGAAGATGACAGCCGCTCACCATTGCTCTACTCCAGGCGCTCAGTCTCCCATGTCTCC 300
DB 447 AAGAAGATGACAGCCGCTCACCATTGCTCTACTCCAGGCGCTCAGTCTCCCATGTCTCC 506
QY 301 ATGTAGAGTGCAGCTTCTCTCCCGAGTACTTCTCAGCTCCGGGCTGGCTCAGGACCC 360
DB 507 ATGTAGAGTGCAGCTTCTCTCCCGAGTACTTCTCAGCTCCGGGCTGGCTCAGGACCC 566
QY 361 GTGTCTCAGTGGCCAGGAACGTTATGAAGCATCAGACCTTAACCTGGAGAGAGGAG 420
DB 567 GTGTCTCAGTGGCCAGGAACGTTATGAAGCATCAGACCTTAACCTGGAGAGAGGAG 593
QY 421 GAAGAAGAGGGGAGAGAGAGGAAGAAGAGAGATGATGAGATGAGATGACAGAT 480
DB 594 ----- 593
QY 481 ATATCTCTGAGAGCAAAAGCCCTGTCAAAACAAGTCAAAAAGGCTGGTCCCGAGAAGCAG 540
DB 594 ----- 593
QY 541 GCGAGCGTGGCTAAGAAAAAAGCTGAAAAAAGAAGAGAGAAATAAGAGCCAGCGTT 600
DB 594 -----GAAAAAAAGCTGAAAAAAGAAGAAGAGAAATAAGAGCCAGCGTT 639
QY 601 AGAGACAAGAGCCCTGTGAAAAAAGGCCAAAGCCACAGCCAGAGCCAAAGCCAGATTC 660
DB 640 AGAGACAAGAGCCCTGTG-AAAAAGGCCAAAGCCACAGCCAGAGCCAAAGCCAGATTC 698
QY 661 AAGAATGAGAGCCAGCCCTTGGGGGCGACGCTGCAAAAGTGGGCTTCCCTGGGCTGTG 720
DB 699 AAGAATGAGAGCCAGCCCTTGGGGGCGACGCTGCAAAAGTGGGCTTCCCTGGGCTGTG 758
QY 721 CTGCAGGACAGGGTGGCCCTGTCCAGCCCTTCCACCTGTGTCTGAATGCAACAGGGT- 779
DB 759 CTGCAGGACAGGGTGGCCCTGTCCAGCCCTTCCACCTGTGTCTGAATGCAACNGGGNT 818
QY 780 --GTTGGGGGGAACATG-AGAGCCCTCACCCTCACT 816
DB 819 GGTTCGCGGGGACACCATGAAGAGCCCTCACCCTCACT 858

RESULT 8
BQ878312 872 bp mRNA linear EST 16-AUG-2002
LOCUS BQ878312
DEFINITION AGENCOURT_8072779 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6087652
5' mRNA sequence.
ACCESSION BQ878312
VERSION BQ878312.1 GI:22270320
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
1 (bases 1 to 872)
AUTHORS NIH-MGC
TITLE http://mhc.ncl.nih.gov/
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
Contact: Robert Strausberg, Ph.D.
Email: cgabs-remail.nih.gov
Tissue Procurement: DCTP/DTP
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:
http://image.lnl.gov
Plate: LCM2324 row: j column: 05
High quality sequence stop: 548.

FEATURES
source
1. .872
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:6087652"
/clone_lib="NIH_MGC_112"
/tissue_type="melanotic melanoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: skin; Vector: pOTB7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC Library."

BASE COUNT 192 a 265 c 259 g 145 t 11 others
ORIGIN

Query Match 43.9%; Score 405.8; DB 14; Length 872;
Best Local Similarity 76.1%; Pred. No. 1.8e-58;
Matches 602; Conservative 0; Mismatches 20; Indels 169; Gaps 2;

QY 1 CAGCCCGCTTCTCTGCGGAGCCATGAAATCTCAGTAGCCGAGTAGCAGGAGGAAAG 60
DB 207 CAGCCCGCTTCTCTGCGGAGCCATGAAATCTCAGTAGCCGAGTAGCAGGAGGAAAG 266
QY 61 GCAGTGACGACCGTGTCTGGGGCTGCGAGCTCAGTCAGAGAGGCGGACTTGACCTTC 120
DB 267 GCAGTGACGACCGTGTCTGGGGCTGCGAGCTCAGTCAGAGAGGCGGACTTGACCTTC 326
QY 121 AGACCCCACTGGAGGGGAGAGAGAGCTGCAGGCTGTTCCTTATACGATTTGCTTGGGG 180
DB 327 AGACCCCACTGGAGGGGAGAGAGAGCTGCAGGCTGTTCCTTATACGATTTGCTTGGGG 386
QY 181 GAGAAAGCCAAAGAGAGATGATTCGCGTGGAGATCCTGCCCCAGCAAAACCAGAGGAC 240
DB 387 GAGAAAGCCAAAGAGAGATGATTCGCGTGGAGATCCTGCCCCAGCAAAACCAGAGGAC 446
QY 241 AAGAAGATGACAGCCGCTCACCATTGCTCTACTCCAGGCGCTCAGTCTCCCATGTCTCC 300
DB 267 AAGAAGATGACAGCCGCTCACCATTGCTCTACTCCAGGCGCTCAGTCTCCCATGTCTCC 506
QY 301 ATGTAGAGTGCAGCTTCTCTCCCGAGTACTTCTCAGCTCCGGGCTGGCTCAGGACCC 360
DB 507 ATGTAGAGTGCAGCTTCTCTCCCGAGTACTTCTCAGCTCCGGGCTGGCTCAGGACCC 566
QY 361 GTGTCTCAGTGGCCAGGAACGTTATGAAGCATCAGACCTTAACCTGGAGAGAGGAG 420
DB 567 GTGTCTCAGTGGCCAGGAACGTTATGAAGCATCAGACCTTAACCTGGAGAGAGGAG 593
QY 421 GAAGAAGAGGGGAGAGAGAGGAAGAAGAGAGATGATGAGATGAGATGAGATGAGAT 480
DB 594 ----- 593
QY 481 ATATCTCTGAGAGCAAAAGCCCTGTCAAAACAAGTCAAAAAGGCTGGTCCCGAGAAGCAG 540


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Db 594 ----- 593
QY 541 GCGAGCGTGGCTAAGAAAAAAGCTGGAAAAAGAGAGAGGAATTAAGAGCCGCTT 600
Db 594 -----GAAAAAAGCTGGAAAAAAGAGAGAGGAATTAAGAGCCGCTT 639
QY 601 AGAGACAAGAGCCCTGTGAAAAAGGCCAAAGCCACAGCCAGAGCCAGAAGCGAGATTTC 660
Db 640 AGAGACAAGAGCCCTGTGAAAAAGGCCAAAGCCACAGCCAGAGCCAGAAGCGAGATTTC 699
QY 661 AAGAATGAGAGAGCCACGCGCTTGGGGGGGCGACGGTGCAGAAAGTGCGCTTCCCTGGGCTGTG 720
Db 700 AAGAATGAGAGAGCCACGCGCTTGGGGGGGCGACGGTGCAGAAAGTTCCTTCCCTGGGCTGTG 759
QY 721 CTGCAGGCAAGGGTGGCCCTGTCCAGCCCGC--TCCACCTGTGTCTGAATGCAACAGGGG 778
Db 760 CTGCAGGCAAGGGTGGCCCTGTCCAGCCCGCCTCCCTGCTGTGAATGCAACAGAGG 819
QY 779 TGTTCGGGGGG 789
Db 820 GGTGTTGGGGG 830

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RESULT 9

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LOCUS BF594409/c 405 bp mRNA linear EST 12-DEC-2000
DEFINITION 7105a05.x1 NCI_CGAP_CO16 Homo sapiens CDNA clone IMAGE:3324560 3'
similar to contains Alu repetitive element; contains element MER28
repetitive element ;, mRNA sequence.

```

ACCESSION

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BF594409
BF594409.1 GI:11686733
EST.

```

KEYWORDS

SOURCE

ORGANISM

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Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```

REFERENCE

```

1 (bases 1 to 405)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)

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JOURNAL

```

Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Ian Kirsch, M.D., Michael R. Emmert-Buck, M.D.,
Ph.D.

```

```

CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL, send email to:
info@image.llnl.gov

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Seq primer: -40UP from Gibco
High quality sequence stop: 404.
Location/Qualifiers
1. 405

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FEATURES

source

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3324560"
/clone_lib="NCI_CGAP_CO16"
/tissue_type="Colon tumor, RER+"
/lab_host="DH10B"
/notes="Organ: colon; Vector: pT7T3D-Pac (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI;
Plasmid DNA from the normalized library NCI_CGAP_CO10 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneIDs 1057416-1061255, and 114584-1145351).
Subtraction by Bento Soares and M. Fatima Bonaldo."

```

BASE COUNT

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54 a 120 c 107 g 124 t
ORIGIN

```

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Query Match 43.8%; Score 405; DB 12; Length 405;
Best Local Similarity 100.0%; Pred. No. 3.2e-58;
Matches 405; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 488 TGGAGGAGCAAGGCCCTGTCAACAAGTCAAAAGGCTGTGCCCCAGAGCGAGCG 547
Db 405 TGGAGGAGCAAGGCCCTGTCAACAAGTCAAAAGGCTGTGCCCCAGAGCGAGCG 346
QY 548 TGGCTAAGAAAAAAGCTGGAAAAAGAGAGGAATTAAGAGCCAGCTTAGAGACA 607
Db 345 TGGCTAAGAAAAAAGCTGGAAAAAGAGAGGAATTAAGAGCCAGCTTAGAGACA 286
QY 608 AGAGCCCTGTGAAAAAAGGCCAAAGCCACAGCCAGAGCCAGAGCCAGATTCAAGAAAT 667
Db 285 AGAGCCCTGTGAAAAAAGGCCAAAGCCACAGCCAGAGCCAGAGCCAGATTCAAGAAAT 226
QY 668 GAGGAGCCAGCGCTTGGGGGGGCGACGGTGCAGAAAGTGCGCTTCCCTGGGCTGTGCTGCAGG 727
Db 225 GAGGAGCCAGCGCTTGGGGGGGCGACGGTGCAGAAAGTGCGCTTCCCTGGGCTGTGCTGCAGG 166
QY 728 CACAGGGTGGCCCTGTCCAGCCCGCTCCACCTGTGTCTGAATGCAACAGGGGTGTCGGGG 787
Db 165 CACAGGGTGGCCCGCTGTCCAGCCCGCTCCACCTGTGTCTGAATGCAACAGGGGTGTCGGGG 106
QY 788 GGCACATGAGAGAGCCCTGCACCCCAACTCTCCACTTTCAGAGAGGCCCCAGTGAAGAGC 847
Db 105 GGCACATGAGAGAGCCCTGCACCCCAACTCTCCACTTTCAGAGAGGCCCCAGTGAAGAGC 46
QY 848 CCCACCTCGGGGTCAATTAAGTTGCTGCTCAGGAAAAA 892
Db 45 CCCACCTCGGGGTCAATTAAGTTGCTGCTCAGGAAAAA 1

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RESULT 10

LOCUS

```

AW731946/c 399 bp mRNA linear EST 21-APR-2000
DEFINITION ba03e06.x1 NIH_MGC_7 Homo sapiens CDNA clone IMAGE:2823298 3', mRNA
sequence.

```

ACCESSION

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AW731946
AW731946.1 GI:7632268
EST.

```

KEYWORDS

SOURCE

ORGANISM

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Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```

REFERENCE

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1 (bases 1 to 399)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)

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JOURNAL

```

Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
CDNA Library Preparation: Ling Hong/Rubin Laboratory CDNA Library
Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
image.llnl.gov/image/html/resources.shtml

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Seq primer: -40UP from Gibco
High quality sequence stop: 381.
Location/Qualifiers
1. 399

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FEATURES

source

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2823298"
/clone_lib="NIH_MGC_7"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: lung; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; CDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Size-selected >500bp for average

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Insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."

BASE COUNT 53 a 116 c 108 g 122 t
ORIGIN

Query Match 43.0%; Score 397.4; DB 10; Length 399;
Best Local Similarity 99.7%; Pred. No. 5.9e-57;
Matches 398; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 493 GAGCAAAAGCCCTGTCAACAAGTCAAAAGGCTGTGCCCCAGAGAGCAGCGCGTGGCT 552
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Db 399 GAGCAAAAGCCCTGTCAACAAGTCAAAAGGCTGTGCCCCAGAGAGCAGCGCGTGGCT 340
QY 553 AAGAAAAAAGCTGGAAGAAGAGGAATAGAGCCAGCGTTAGAGCAAGAGC 612
|||
Db 339 AAGAAAAAAGCTGGAAGAAGAGGAATAGAGCCAGCGTTAGAGCAAGAGC 280
QY 613 CCTGTGAAAAAGGCCAAGCCACAGCCAGCCAGGATTCAGAAATGAGGA 672
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Db 279 CCTGTGAAAAAGGCCAAGCCACAGCCAGGATTCAGAAATGAGGA 220
QY 673 GCCACGCTTGGGGGCGACCGTGCAAAAGTGGGCTTCCCTGGGCTGTGCTGCAGGACAG 732
|||
Db 219 GCCACGCTTGGGGGCGACCGTGCAAAAGTGGGCTTCCCTGGGCTGTGCTGCAGGACAG 160
QY 733 GGTGCCCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCAACAGGGGTGTGCGGGGCCAA 792
|||
Db 159 GGTGCCCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCAACAGGGGTGTGCGGGGCCAA 100
QY 793 CATGAGAGCCCCCTCACCCCACTCTCCACTTTTCAGAGAGGCCCCCAGTGAAGAGCCCCAC 852
|||
Db 99 CATGAGAGCCCCCTCACCCCACTCTCCACTTTTCAGAGAGGCCCCCAGTGAAGAGCCCCAC 40
QY 853 CTCGGGGTCAATAAAGTTGCTGTGAGGAAAAAAA 891
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Db 39 CTCGGGGTCAATAAAGTTGCTGTGAGGAAAAAAA 1

RESULT 11
B0684424 935 bp mRNA linear EST 15-JUL-2002
LOCUS
DEFINITION AGENCOURT_8209062 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6260492
5', mRNA sequence.
ACCESSION B0684424
VERSION B0684424.1 GI:21797103
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS NIH-MGC http://mgi.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: c9apbs-remail.nih.gov
Tissue Procurement: DCTD/DTP
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLCM2419 row: k column: 21
High quality sequence stop: 635.

FEATURES

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:6260492"
/clone_1ib="NIH_MGC_112"

/tissue_type="melanotic melanoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: skin; Vector: pOTB7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC Library."

BASE COUNT 186 a 304 c 269 g 175 t 1 others
ORIGIN

Query Match 42.2%; Score 389.6; DB 14; Length 935;
Best Local Similarity 98.7%; Pred. No. 8.9e-56;
Matches 392; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 CAGCCCGCTTCTCTGCCCCGAGCCATGATCTCAGTAGCGCCAGTAGCAGGAGAAAAAG 60
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Db 207 CAGCCCGCTTCTCTGCCCCGAGCCATGATCTCAGTAGCGCCAGTAGCAGGAGAAAAAG 266
QY 61 GCAGTAGCAGCCGTCTCTGGGGCTGCGAGCTCAGTCAGAGAGCGGACTTGACCTTC 120
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Db 267 GCAGTAGCAGCCGTCTCTGGGGCTGCGAGCTCAGTCAGAGAGCGGACTTGACCTTC 326
QY 121 AGACCCAGCTGAGGGGAGAGCAGAGCTGCAGGCTGTGCTTATACGATTGCTGGGG 180
|||
Db 327 AGACCCAGCTGAGGGGAGAGCAGAGCTGCAGGCTGTGCTTATACGATTGCTGGGG 386
QY 181 GAGAAAGCCAAAGAGAGATGATCGCGGTGAGATCCTGCCCCCAGCAAAACAGAGAGAC 240
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Db 387 GAGAAAGCCAAAGAGAGATGATCGCGGTGAGATCCTGCCCCCAGCAAAACAGAGAGAC 446
QY 241 AAGAAGATGACCGGTCACCATTTGCTCCTCAGGCTCAGTCCCTCCATGCTCC 300
|||
Db 447 AAGAAGATGACCGGTCACCATTTGCTCCTCAGGCTCAGTCCCTCCATGCTCC 506
QY 301 ATGGTAGAGTGCAGCTTCTCCCCAGTTACTTTCCAGCTCCGGGCTGCTCAGGACCC 360
|||
Db 507 ATGGTAGAGTGCAGCTTCTCCCCAGTTACTTTCCAGCTCCGGGCTGCTCAGGACCC 566
QY 361 GTGTTCTCAGTGGCCAGGAACGTTATGAAGCATCAG 397
|||
Db 567 GTGTTCTCAGTGGCCAGGAACGTTATGAAGCATCAG 603

RESULT 12
B0682257 966 bp mRNA linear EST 15-JUL-2002
LOCUS
DEFINITION AGENCOURT_8194971 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6259047
5', mRNA sequence.
ACCESSION B0682257
VERSION B0682257.1 GI:21794936
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS NIH-MGC http://mgi.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: c9apbs-remail.nih.gov
Tissue Procurement: DCTD/DTP
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLCM2415 row: o column: 16
High quality sequence stop: 518.

FEATURES
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1. 966

/organism="Homo sapiens"
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/tissue_type="melanotic melanoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: skin; Vector: pOTB7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC library."

BASE COUNT 205 a 300 c 295 g 159 t 7 others
ORIGIN

Query Match 41.0%; Score 378.4; DB 14; Length 966;
Best Local Similarity 74.0%; Pred. No. 6.5e-54;
Matches 668; Conservative 0; Mismatches 56; Indels 179; Gaps 8;

QY 1 CAGCCCGCTTCTCTGCGCGGAGCCATGATCTCAGTAGCGCCAGTAGCAGGAGGAAAG 60
DB 207 CAGCCCGCTTCTCTGCGCGGAGCCATGATCTCAGTAGCGCCAGTAGCAGGAGGAAAG 266
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DB 267 GCAGTGACGACCGCTGCTGGGGCTGCGAGCTCAGTCAGGAGGCGGACTTGACCTTC 326
QY 121 AGACCCAGCTGAGGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 180
DB 327 AGACCCAGCTGAGGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 386
QY 181 GAGAAAGCCAAAGAGAGATGC-ATCGCTGAGAGATCCTGCCCCAGCAAAACAGAGAGA 239
DB 387 GAGAAAGCCAAAGAGAGATGCATCGCTGAGAGATCCTGCCCCAGCAAAACAGAGAGA 446
QY 240 CAAGAAGATGACGCGGCTCACCATTGCTCAGGCTCAGTCTCCCATGCTCTC 299
DB 447 CAAGAAGATGACGCGGCTCACCATTGCTCAGGCTCAGTCTCCCATGCTCTC 506
QY 300 CATGG--TAGAGTGACAGCTTCTCCCGAGTTACTTTCAGCTCCGGCTGGCTCAGG- 356
DB 507 CATGGTGAGGAGTGCAGCTTCTCCCGAGTTACTTTCAGCTCCGGCTGGCTCAGG 566
QY 357 -ACCCGCTTCTCAGTGGCCAGAGAGTTATGAGCATCAGACCTAACCTGGGAGAGG 415
DB 567 ACCCGTGTCTCTCAGTGGCCAGAGAGTTAT----- 598
QY 416 AGAGAGAGAGAGAGGAGGAGGAGAGAGAGAGAGAGAGATGATGAGATGAGATG 475
DB 599 ----- 598
QY 476 CAGATATATCTCTGAGAGAGCAAGCCCTGTCAAAACAGTCAAAAGGCTGTGCCAGAG 535
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DB 599 -----GAAAAAAGCTGAAAAAAGAGAGAGAAATAGAGAGCA 639
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DB 640 GCGTTAGAGACAGAGCCCTGTGAAAAAGGCCAAAGCCAGCCAGAGAGCAAGAGAGCCAG 699
QY 656 GATTCAAGAAATGAGAGAGCCAC--GCCTTGGGGGAGGAGGAGGAGGAGGAGGAGGAG 712
DB 700 GATTCAAGAAATGAGAGAGCCCGCTTGGGGGAGGAGGAGGAGGAGGAGGAGGAGGAG 759
QY 713 GGGCTGTCTGACGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 772
DB 713 GGGCTGTCTGACGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 772

DB 760 GCGCTGTGCTGACAGGAGCAGGGTGCCCTGCGCCAGCCCCCTCACCTGGGCTTGAATGCA 819
QY 773 CAGGGGTG--TTGGGGGGCAACATGAGAGCCCTCACCCTCAACTCTCCACTTTACAGA 830
DB 820 CCGGGGGGTTGCGGGGNCACCTGAGAACCCCTCCCTCCCACTCTCCANTTTTCGGG 879
QY 831 --GGCCCCAGTGAAGAGCCCTCAGGCTGACATTAAGTTCCCTGTCAGGAGAAA 888
DB 880 AGGGCCCCAGNGCAAAACCCCTTGGGGTGAAAAAACCCGTGCTGGGCGGAGNAC 939
QY 889 AAA 891
DB 940 AAA 942

RESULT 13
AW002370/c 361 bp mRNA linear EST 09-MAR-2000
LOCUS

DEFINITION wu61a07.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:2524500 3',
mRNA sequence.

ACCESSION AW002370
VERSION AW002370.1 GI:5849286

KEYWORDS EST.
SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.

cdNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/btrp/image/image.html
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Seq primer: -40UP from Gibco.

FEATURES
source

Location/Qualifiers
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/clone="IMAGE:2524500"
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/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA
from the normalized library NCI_CGAP_GC4 was prepared, and
ss circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (clonids
1257096-1258631, 1469064-1470983, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 50 a 107 c 93 g 111 t
ORIGIN

Query Match 38.7%; Score 357.8; DB 10; Length 361;
Best Local Similarity 99.4%; Pred. No. 2.5e-50;
Matches 359; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 533 AGAAGCAGGCGAGCTGCTAAGAAAAAAGCTGAAAAAAGAGAGAGAAATAGAG 592
DB 361 AGAAGCAGGCGAGCTGCTAAGAAAAAAGCTGAAAAAAGAGAGAGAAATAGAG 302

QY 593 CCAGCGTTAGAGACAAGACCCCTGTGAAAAAGCCAAAGCCACAGCCAGAGCCCAAGAAGC 652
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Db 301 CCAGCGTTAGAGACAAGACCCCTGTGAAAAAGCCAAAGCCACAGCCAGAGCCCAAGAAGC 242
QY 653 CAGGATTCAGAAATGAGAGCCACGCGCTTGGGGGGCACGGTGCAGAACTGGGCCCTCCCT 712
|||||
Db 241 CAGGATTCAGAAATGAGAGCCACGCGCTTGGGGGGCACGGTGCAGAACTGGGCCCTCCCT 182
QY 713 GGGCTGTGCTGCAGGACAGGGTGGCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCCAA 772
|||||
Db 181 GGGCTGTGCTGCAGGACAGGGTGGCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCCAA 122
QY 773 CAGGCGTGTGCGGGGGCAACATGAGAGCCCTCAACCCCAACTCTCCACTTTCAGGAGG 832
|||||
Db 121 CAGGCGTGTGCGGGGGCAACATGAGAGCCCTCAACCCCAACTTTCAGACTTTCAGGAGG 62
QY 833 CCCCCAGTGAAGAGCCCCCAGCTCGGGGTCAACAATAAAGTTGCTGTCAGGAAAAA 892
|||||
Db 61 CCCCCAGTGAAGAGCCCCCAGCTCGGGGTCAACAATAAAGTTGCTGTCAGGAAAAA 2
QY 893 A 893
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Db 1 A 1
RESULT 14
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LOCUS 7394b04.x1 NCI_CGAP_GC6 Homo sapiens CDNA clone IMAGE:3442494 3',
DEFINITION mRNA sequence.
ACCESSION BF592761
VERSION BF592761.1 GI:11685085
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 361)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLES National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL, send email to:
info@image.llnl.gov
Seq primer: -40UP from Gibco.
FEATURES
source
1. 361
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3442494"
/clone_lib="NCI_CGAP_GC6"
/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA
from the normalized library NCI_CGAP_GC4 was prepared, and
ss circles were made in vitro. Following HAP purification,
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (clones
1257096-1258631, 1469064-1470983, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 50 a 106 c 94 g 111 t
ORIGIN

Query Match 38.7%; Score 357.8; DB 12; Length 361;
Best Local Similarity 99.4%; Pred. No. 2.5e-50;
Matches 359; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 361 AGAAGCAGGCGCGCTGGCTAAGAAAAAAGCTGGAAGAAAGAGAGAGAAATAAGAG 302
QY 593 CCAGCGTTAGAGACAAGACCCCTGTGAAAAAGCCAAAGCCACAGCCAGAGCCCAAGAAGC 652
|||||
Db 301 CCAGCGTTAGAGACAAGACCCCTGTGAAAAAGCCAAAGCCACAGCCAGAGCCCAAGAAGC 242
QY 653 CAGGATTCAGAAATGAGAGCCACGCGCTTGGGGGGCACGGTGCAGAACTGGGCCCTCCCT 712
|||||
Db 241 CAGGATTCAGAAATGAGAGCCACGCGCTTGGGGGGCACGGTGCAGAACTGGGCCCTCCCT 182
QY 713 GGGCTGTGCTGCAGGACAGGGTGGCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCCAA 772
|||||
Db 181 GGGCTGTGCTGCAGGACAGGGTGGCCCTGTCCAGCCCTCCACCTGTGTCTGAATGCCAA 122
QY 773 CAGGCGTGTGCGGGGGCAACATGAGAGCCCTCAACCCCAACTCTCCACTTTCAGGAGG 832
|||||
Db 121 CAGGCGTGTGCGGGGGCAACATGAGAGCCCTCAACCCCAACTTTCAGACTTTCAGGAGG 62
QY 833 CCCCCAGTGAAGAGCCCCCAGCTCGGGGTCAACAATAAAGTTGCTGTCAGGAAAAA 892
|||||
Db 61 CCCCCAGTGAAGAGCCCCCAGCTCGGGGTCAACAATAAAGTTGCTGTCAGGAAAAA 2
QY 893 A 893
|
Db 1 A 1
RESULT 15
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DEFINITION mRNA sequence.
ACCESSION BF057162
VERSION BF057162.1 GI:10811058
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 361)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLES National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs-r@mail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL, send email to:
info@image.llnl.gov
Seq primer: -40UP from Gibco.
FEATURES
source
1. 361
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3444661"
/clone_lib="NCI_CGAP_GC6"
/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA

from the normalized library NCI CGAP_G4 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clones 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 50 a 107 c 93 g 111 t
ORIGIN

Query Match 38.7%; Score 357.8; DB 12; Length 361;
Best Local Similarity 99.4%; Pred. No. 2.5e-50;
Matches 359; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 533 AGAAGCAGGCGAGCGCTGGCTAAGAAAAAGCTGGAAGAGAGAGGAATAGAG 592
Db 361 AGAAGCAGGCGAGCGCTGGCTAAGAAAAAGCTGGAAGAGAGAGGAATAGAG 302
QY 593 CCAGCGTTAGAGACAGAGCCCTGTGAAAAAGGCCAAGCCACAGCCAGAGCCAGAGAGC 652
Db 301 CCAGCGTTAGAGACAGAGCCCTGTGAAAAAGGCCAAGCCACAGCCAGAGCCAGAGAGC 242
QY 653 CAGCATTCAGAAATGAGGACGCGCTTGGGGGGCAGCGTGCAAAAGTGCGCTTCCCT 712
Db 241 CAGCATTCAGAAATGAGGACGCGCTTGGGGGGCAGCGTGCAAAAGTGCGCTTCCCT 182
QY 713 GGGCTGTGCTGACAGGACAGCGTGCCCTGTCCAGCCCTCCACCTGTGTGAATGCAA 772
Db 181 GGGCTGTGCTGACAGGACAGCGTGCCCTGTCCAGCCCTCCACCTGTGTGAATGCAA 122
QY 773 CAGGGGTGTGCGGGGCAACATGAGAGCCCTCAACCCCAACTCTCCACTTTCAGAGG 832
Db 121 CAGGGGTGTGCGGGGCAACATGAGAGCCCTCAACCCCAACTCTCCACTTTCAGAGG 62
QY 833 CCCCCAGTGAAGAGCCCACTCGGGTCACAATTAAGTTGCTGTGAGGAAAAAAA 892
Db 61 CCCCCAGTGAAGAGCCCACTCGGGTCACAATTAAGTTGCTGTGAGGAAAAAAA 2
QY 893 A 893
Db 1 A 1

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Job time : 2248 secs

